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<p>(54) Title: MATERIALS AND METHODS FOR THE MODIFICATION OF ISOPRENOID CONTENT, COMPOSITION AND METABOLISM</p>		
<p>(57) Abstract</p> <p>Novel isolated polynucleotides associated with plant isoprenoid biosynthetic pathways are provided, together with genetic constructs comprising such sequences. Methods for the modulation of the content, structure and metabolism of polypeptides involved in an isoprenoid biosynthetic pathway in target organisms are also disclosed, the methods comprising incorporating one or more of the polynucleotides or genetic constructs of the present invention into the genome of a target organism. Modulation of the content, structure and metabolism of such polypeptides produces modifications in the content, structure and metabolism of isoprenoids in the target organism.</p>		

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MATERIALS AND METHODS FOR THE MODIFICATION OF ISOPRENOID CONTENT, COMPOSITION AND METABOLISM

5

Technical Field of the Invention

This invention relates materials and methods for modifying the content, composition and metabolism of isoprenoids in plants and other organisms. More particularly, this invention relates to polypeptides involved in the synthesis of isoprenoid compounds, such as terpenoid and steroid compounds, polynucleotides encoding such polypeptides, expression of such polypeptides, and methods for modulating the composition and/or expression levels of such polypeptides, thereby modulating isoprenoid content, composition, and metabolism.

15 Background of the Invention

Isoprenoids form a large family of naturally occurring compounds, with over 20,000 distinct compounds having been described. The isoprenoids include vitamins A, D, E, and K, first recognized as fatty materials essential to the normal growth of animals, and numerous biological pigments. In plants, isoprenoid compounds, including terpenoid and steroid compounds, include hormones such as gibberellic acid and abscisic acid, pigments, electron carriers, membrane components (phytosterols), phytotoxins, antibiotics, flavors such as menthol, vitamins, macromolecular compounds such as rubber and guttapercha, and others.

Isoprene compounds, or prenyl lipids, are composed of one or more basic isoprene skeleton(s) (C_5) formed by the decarboxylation of mevalonate-5-pyrophosphate. From the isopentenyl pyrophosphate ("active isoprene" or "IPP") and the isomeric dimethylallyl pyrophosphate, the geranyl pyrophosphate (C_{10}) may be formed by "head-tail" condensation. By linkage of a further C_5 unit, farnesyl pyrophosphate (C_{15}) is formed. Further extension by "head-tail" or "tail-tail" condensation leads to C_{20} , C_{30} and C_{40} compounds, as well as the higher molecular terpenoids. A schematic diagram of the basic biosynthetic pathways of isoprene compounds is shown in Fig. 1.

IPP is the branching point for a large variety of biologically significant molecules, including isoprenoids, carotenoids, and various sterols in different eukaryotic organisms

(mycosterols, phytosterols and zoosterols). In animals, cholesterol are precursors for several hormones and bile acids. Fungal ergosterol and mammalian cholesterol arise from IPP via squalene oxide and lanosterol, while higher plant sterols, like campesterol and sitosterol, are produced by cyclization of squalene oxide to cycloartenol and by further
5 plant-specific enzymes.

Plant cells contain an intriguing diversity of a subclass of isoprenoids called terpenoids, most of which are cyclic with one or more rings. Terpenes in plants are divided into several classes, including sesquiterpenes, mono-, di-, triterpenes, etc. (Bohlmann *et al.*, *Proc. Natl. Acad. Sci. USA* 95:4126-4133, 1998). Terpenoids are formed by linking
10 isoprene units (C_5H_8) synthesized from acetate. Terpenoids include isoprene (C_5H_8) compounds, including isopen-tenylpyrophosphate and active isoprene; monoterpene ($C_{10}H_{16}$) compounds, including geraniol, and from which menthol, camphor, pinene and citronellal are derived; sesquiterpene ($C_{15}H_{24}$) compounds, including farnesol, from which zingiberene, ubiquinone, plastoquinone, abscisic acid and rishitine are derived; diterpene
15 ($C_{20}H_{32}$) compounds, such as geranylgeraniol, from which phytol, kaurene, gibberellin acid and fusicoccin are derived; triterpene ($C_{30}H_{48}$) compounds, including squalene, from which steroids and saponins are derived; tetraterpene ($C_{40}H_{64}$) compounds, including phytoene and carotenes; and polyterpene (C_5H_8)_n compounds, including rubber and guttapercha.

20 Synthase enzymes producing terpenes are thought to be of common evolutionary origin, lacking close similarity to other enzymes (except prenyltransferases). Most synthase enzymes have the ability to produce a variety of end-products from a single substrate. This may explain, in part, the enormous diversity of terpenoid compounds found in plants (Mitchell-Olds *et al.*, *Trends in Plant Science* 3(9):362-365, 1998). Complex
25 terpene mixtures are thought to be important plant defensive compounds, their diversity and synergistic action delaying development of resistance in herbivores and pathogens (Langenheim J, *J. Chem. Ecol.* 20:1223-1280, 1994).

Plant terpenoids also have many known medicinal effects, and some plant isoprenoid compounds are administered as drugs. Taxol, which has proven to be
30 efficacious in treating cancer, for example, is derived from terpenoid compounds. Dietary isoprenoids have been suggested to suppress mevalonate pathway, thereby affecting cancer and cardiovascular disease (Elson CE, *J. Nutr.* 125(6 Suppl):1666S-1672S, 1995). Farnesol, the last precursor common to all branches of the mevalonate pathway, has been

demonstrated to inhibit calcium channels in muscle cells (Roulette J-B, *J. Biol. Chem.* 51:32240-32246, 1997).

Ubiquinone and plastoquinone, which are also isoprenoid derivatives, function as electron carriers in the production of ATP in mitochondria and chloroplasts. In most mammalian tissues, ubiquinone (also called coenzyme Q) has ten isoprene units. Plastoquinone is the plant equivalent of ubiquinone. In their role as electron carriers, both ubiquinone and plastoquinone can accept either one or two electrons and either one or two protons to be reduced.

A remarkable role for isoprenyl intermediates has recently been discovered in studies of a protein that is implicated in human cancers and is known to associate with membranes through a covalently bound isoprenyl lipid. This protein, the RAS PROTEIN, is the product of the gene, a mutant version of a normal protein and a number of related GTP-binding proteins. The normal protein and the number of related GTP-binding proteins are known to act in signal transductions triggered by neurotransmitters, hormones, growth factors and other extracellular signals.

Quantitative and qualitative modifications in plant terpenoid content are known to be induced by external factors such as herbivore attack and wounding (Bohlmann J *et al.*, *Proc. Natl. Acad. Sci. USA* 95:6756-6761, 1998). Synthesis of cell terpenoids can also be induced by infection with pathogens. Even agricultural pest insects can be repelled by pine oil terpene compounds: monoterpenes carene, limonene and cymene deter onion flies (Ntiamoah Ya, *Entom. Exp. et Appl.* 79:219-226, 1996).

While the chemical diversity of isoprenoids is well known, and many of the metabolic pathways have been tentatively identified, few of the genes encoding enzymes responsible for the synthesis of isoprenoid compounds have been identified. The present invention is therefore directed to providing novel polynucleotides encoding polypeptides involved in the biosynthesis of isoprenoids, and providing methods for modifying the expression and composition of such polypeptides, thereby modulating isoprenoid content, composition, and metabolism.

Sequencing of the genomes, or portions of the genomes, of numerous biological materials, including humans, animals, microorganisms and various plant varieties, has been and is being carried out on a large scale. Polynucleotides identified using sequencing techniques may be partial or full-length genes, and may contain open reading frames, or portions of open reading frames, that encode polypeptides. Putative polypeptides may be

determined based on polynucleotide sequences. The sequencing data relating to polynucleotides thus represents valuable and useful information.

Polynucleotides may be analyzed for novelty by comparing identified sequences to sequences published in various public domain databases, such as EMBL. Newly
5 identified polynucleotides and putative polypeptides may also be compared to polynucleotides and polypeptides contained in databases to ascertain homology to known polynucleotides and polypeptides. In this way, the degree of similarity or identity or homology of polynucleotides and polypeptides having an unknown function may be determined relative to polynucleotides and polypeptides having known functions.

10 U.S. Patent 5,589,619 discloses materials and methods for increasing squalene and sterol accumulation in higher plants by modifying the copy number of a gene encoding a polypeptide having HMG-CoA reductase activity. Genetic materials, including polynucleotides, polypeptides, DNA molecules, and the like, relating to HMG-CoA reductase activity are disclosed, as well as methods for transforming plant cells and
15 producing transgenic plants.

U.S. Patent 5,689,047 discloses stilbene synthase genes derived from grapevines, as well as the use of those genes in vectors and transformed microorganisms, as well as transformed plant cells and plants.

U.S. Patent 5,753,507 discloses plant polynucleotides encoding geraniol/nerol 10 –
20 hydroxylase ($G_{10}H$), as well as methods for using complete and partial polynucleotides as probes, and methods for expressing $G_{10}H$ and enhancing levels of terpenoid indole alkaloid and iivoid insect pheromone produced by a plant.

The following U.S. Patents disclose isoprenoid compounds or related compounds, or methods for using such compounds: U.S. Patent 5,429,939; U.S. Patent 5,444,166; U.S.
25 Patent 5,460,949; U.S. Patent 5,470,832; U.S. Patent 5,474,925; U.S. Patent 5,495,070; U.S. Patent 5,521,078; U.S. Patent 5,545,816; U.S. Patent 5,547,856; U.S. Patent 5,569,832; U.S. Patent 5,580,963; U.S. Patent 5,597,718; U.S. Patent 5,670,349; U.S. Patent 5,674,485; U.S. Patent 5,684,238; U.S. Patent 5,689,056; U.S. Patent 5,691,147; U.S. Patent 5,693,476; and U.S. Patent 5,443,978. The U.S. Patents cited above are
30 incorporated by reference herein in their entireties.

Summary of the Invention

Briefly, the present invention provides isolated polynucleotides encoding polypeptides involved in the production and modification of isoprenoids. Genetic constructs comprising such sequences and methods for the use of such genetic constructs
5 are also provided, together with transgenic cells and plants incorporating such genetic constructs and exhibiting modified isoprenoid content, composition, and metabolism.

In a first aspect, the present invention provides isolated polynucleotide sequences identified in the attached Sequence Listing as SEQ ID NOS: 1-53 and 78-164, variants of those sequences, extended sequences comprising the sequences set out in SEQ ID NOS:
10 1-53, 78-164 and their variants, probes and primers corresponding to the sequences set out in SEQ ID NOS: 1-53, 78-164 and their variants, polynucleotides comprising at least a specified number of contiguous residues of any of the polynucleotides identified as SEQ ID NOS: 1-53 and 78-164 (x-mers), and extended sequences comprising portions of the sequences set out in SEQ ID NOS: 1-53 and 78-164, all of which are referred to herein,
15 collectively, as "polynucleotides of the present invention."

The present invention also provides isolated polypeptide sequences identified in the attached Sequence Listing as SEQ ID NOS: 165-304, polypeptide variants of those sequences, polypeptides comprising the isolated polypeptide sequences and variants of those sequences, polypeptides comprising at least a specified number of contiguous
20 residues of any of the polypeptides identified as SEQ ID NOS: 165-304; and polypeptides comprising portions of the sequences set out in SEQ ID NOS: 165-304.

The polynucleotide sequences identified as SEQ ID NOS: 1-53 and 78-164 were derived from plant sources, namely from *Eucalyptus grandis* and *Pinus radiata*. The polynucleotides of the present invention are primarily "partial" sequences, in that they do
25 not represent a full length gene encoding a full length polypeptide. Such partial sequences may be extended by analyzing and sequencing various DNA libraries using primers and/or probes and well known hybridization and/or PCR techniques. The partial sequences identified as SEQ ID NOS: 1-53 and 78-164 may thus be extended until an open reading frame encoding a polypeptide, a full length polynucleotide and/or gene capable of
30 expressing a polypeptide, or another useful portion of the genome is identified. Such extended sequences, including full length polynucleotides and genes, are described as "corresponding to" a sequence identified as one of the sequences of SEQ ID NOS: 1-53 and 78-164 or a variant thereof, or a portion of one of the sequences of SEQ ID NOS: 1-53

and 78-164 or a variant thereof, when the extended polynucleotide comprises an identified sequence or its variant, or an identified contiguous portion (x-mer) of one of the sequences of SEQ ID NOS: 1-53 and 78-164 or a variant thereof. Similarly, RNA sequences, reverse sequences, complementary sequences, anti-sense sequences, and the like, corresponding to
5 the polynucleotides of the present invention, may be routinely ascertained and obtained using the cDNA sequences identified as SEQ ID NOS: 1-53 and 78-164.

The polynucleotides identified as SEQ ID NOS: 1-53 and 78-164 may contain open reading frames ("ORFs") or partial open reading frames encoding polypeptides. Additionally, open reading frames encoding polypeptides may be identified in extended or
10 full length sequences corresponding to the sequences set out as SEQ ID NOS: 1-53 and 78-164. Open reading frames may be identified using techniques that are well known in the art. These techniques include, for example, analysis for the location of known start and stop codons, most likely reading frame identification based on codon frequencies, etc. Suitable tools and software for ORF analysis are available, for example, on the Internet at
15 <http://www.ncbi.nlm.nih.gov/gorf/gorf.html>. Open reading frames and portions of open reading frames may be identified in the polynucleotides of the present invention. Once a partial open reading frame is identified, the polynucleotide may be extended in the area of the partial open reading frame using techniques that are well known in the art until the polynucleotide for the full open reading frame is identified. Thus, polynucleotides and
20 open reading frames encoding polypeptides may be identified using the polynucleotides of the present invention.

Once open reading frames are identified in the polynucleotides of the present invention, the open reading frames may be isolated and/or synthesized. Expressible DNA constructs comprising the open reading frames and suitable promoters, initiators,
25 terminators, etc., which are well known in the art, may then be constructed. Such DNA constructs may be introduced into a host cell to express the polypeptide encoded by the open reading frame. Suitable host cells may include various prokaryotic and eukaryotic cells, including plant cells.

Polypeptides encoded by the polynucleotides of the present invention may be
30 expressed and used in various assays to determine their biological activity. Such polypeptides may be used to raise antibodies, to isolate corresponding interacting proteins or other compounds, and to quantitatively determine levels of interacting proteins or other compounds.

The present invention also contemplates methods for modulating the polynucleotide and/or polypeptide content and composition of an organism, such methods involving, according to one embodiment, stably incorporating into the genome of the organism a genetic construct comprising one or more polynucleotides of the present invention. In one embodiment, the target organism is a plant, preferably a woody plant, more preferably a woody plant of the *Pinus* or *Eucalyptus* species, and most preferably *Eucalyptus grandis* or *Pinus radiata*. In a related aspect, a method for producing an organism having an altered genotype or phenotype is provided, the method comprising transforming a host cell with a genetic construct of the present invention to provide a transgenic cell, and cultivating the transgenic cell under conditions conducive to growth and regeneration. Organisms having an altered genotype or phenotype as a result of modulation of the level or content of a polynucleotide or polypeptide of the present invention compared to a wild-type organism, as well as components (seeds, etc.) of such organisms and progeny of such organisms, are contemplated by and encompassed within the present invention.

The isolated polynucleotides of the present invention have utility in genome mapping, in physical mapping, and in positional cloning of genes. Additionally, the polynucleotide sequences identified as SEQ ID NOS: 1-53, 78-164, and their variants, may be used to design oligonucleotide probes and primers. Oligonucleotide probes and primers have sequences that are substantially complementary to the polynucleotide of interest over a certain portion of the polynucleotide. Oligonucleotide probes designed using the polynucleotides of the present invention may be used to detect the presence and examine the expression patterns of genes in any organism having sufficiently similar DNA and RNA sequences in their cells using techniques that are well known in the art, such as slot blot DNA hybridization techniques. Oligonucleotide primers designed using the polynucleotides of the present invention may be used for PCR amplifications. Oligonucleotide probes and primers designed using the polynucleotides of the present invention may also be used in connection with various microarray technologies, including the microarray technology used by Synteni (Palo Alto, CA).

The polynucleotides of the present invention may also be used to tag or identify an organism or reproductive material therefrom. Such tagging may be accomplished, for example, by stably introducing a non-disruptive non-functional heterologous

polynucleotide identifier into an organism, the polynucleotide comprising one of the polynucleotides of the present invention.

The polynucleotides of the present invention encode polypeptides that have activity in an isoprenoid biosynthetic pathway. The isoprenoid metabolism-related polynucleotides were isolated from pine and eucalyptus, and putatively identified by DNA and protein similarity searches. Various isoprenoid compounds are well characterized and have useful properties. Methods of the present invention relating to modulating the polynucleotide and/or polypeptide content and composition of an organism and, thereby, modulating the isoprenoid content, composition and metabolism of an organism, are applicable to a wide range of activities. The novel materials and methods of the present invention have a multitude of potential uses: in forestry and agriculture for manipulation of isoprenoid metabolism; in medicine for therapeutic effects, including direct application in diseased organisms or indirect application by transgenic organisms; in fermentation and chemical processing industries involving isoprenoids; and in numerous other applications, some of which are described in the references cited above. In plant applications, manipulating isoprenoid pathways or isoprenoid composition may, for example, affect plant development, pest resistance, and the value of extractives (pinene, myrcene, etc.). In foodstuffs, various isoprenoids affect the nutritional quality and pharmacological properties of the ingested material, e.g. cholesterol or phytosterol composition of animal-derived and plant-derived foods for human or animal consumption. Additionally, isoprenoid pathways control the production of vitamins A, E, and K; plant pigments such as carotene and the phytol chain of chlorophyll; natural rubber; many essential oils, such as the fragrant principles of lemon oil, eucalyptus, and musk; insect juvenile hormone, which controls metamorphosis; dolichols, which serve as lipid-soluble carriers in complex polysaccharide synthesis; and ubiquinone and plastoquinone, electron carriers in mitochondria and chloroplasts. The ubiquitous and varied roles of isoprenoids thus make these compounds and the polynucleotides encoding them attractive targets for biotechnical applications in a variety of fields.

Briefly, the present invention provides isolated polynucleotides encoding polypeptides involved in the synthesis of isoprenoids. The polynucleotides and polypeptides of the present invention have demonstrated similarity to polypeptides that are known to be involved in the synthesis of isoprenoids as shown below in Table 1.

TABLE 1

POLYNUCLEOTIDE SEQ ID NO	POLYPEPTIDE SEQ. ID	POLYPEPTIDE IDENTITY
1	252	Acetylcholinesterase Precursor
2	253	Deoxyxylulosephosphate Synthase (DXPS)
3, 4, 44	254,255,295	Geranyltranstransferase
5, 6	256,266	Farnesyltranstransferase
7, 154	258 241	Squalene Synthetase
8-10, 155-157	259-261 242-244	Squalene Monooxygenase
11	262	Geranylgeranyl-Diphosphate Geranylgeranyltransferase
12	263	Trichodiene Synthase
13, 25, 84-88, 95 115-118	264,276 171-175, 182, 202-205	Pinene Synthase
14, 89, 90	265 176, 177	Abietadine Synthase
15, 91-94, 96-98, 131-135	266 178-181, 183-185, 218-222	Hydroxymethylglutaryl-Coa Reductase (NADPH)
16, 17, 18, 99-102	267,268,269, 186-189	Myrcene Synthase
19, 20, 103, 107, 108	270,271 190, 194, 195	Limonene Synthase
21-23, 109-111	272-274 196-198	Cadinene Synthase
24, 114	275 201	Bisabolene Synthase
26, 27	277,278	Pinene/Myrcene/Limonene Synthase
28, 119-122	279 206-209	Cycloartenol Synthase
29, 124-126	280 211-213	Obtusifoliol Demethylase
30	281	Lupeol Synthase
31, 158, 159	282 245, 246	Udp-Glucose:Sterol Glucosyltransferase
32	283	Hydroxymethylglutaryl-CoA Reductase (NADPH)
33, 34, 160-162	284,285 247-249	Sterolmethyltransferase
35, 136	286 223	Lecithin:Cholesterol Acyl Transferase
36, 137	287 224	Sterol Delta-7 Reductase
37, 38, 138-140	288,289 225-227	Methyl Sterol Oxidase
39	290	Deoxyxylulosephosphate Synthase (DXPS)
40	291	Phosphomevalonate Kinase
41, 50, 141, 142, 146	292,301 228, 229, 233	Diphosphomevalonate Decarboxylase
42, 43, 143	293,294 230	Isopentenyl-Diphosphate Delta- Isomerase

POLYNUCLEOTIDE SEQ ID NO	POLYPEPTIDE SEQ. ID	POLYPEPTIDE IDENTITY
45	296	Estradiol Dehydrogenase
46-49, 144, 145	297-300 231, 232	Furostanol Glucosidase
51, 52, 147-153	302,303 234-240	Oxysterol-Binding Protein
53	304	Sterol Carrier Protein
78, 79, 127-130	165, 166, 214-217	Sterol 14-demethylase
81	168	Sesquiterpene cyclase
82, 83	169, 170	Geranylgeranyl diphosphate
104-106, 164	191-193, 251	CXPS/transketolase
112, 113	199, 200	Sabinene synthase
123	210	Beta-amyrin synthase
163	250	Sterol desaturase

In one embodiment, the isolated polynucleotides comprise a sequence selected from the group consisting of: (a) sequences recited in SEQ ID NOS: 1-53 and 78-164;
 5 (b) complements of the sequences recited in SEQ ID NOS: 1-53 and 78-164; (c) reverse complements of the sequences recited in SEQ ID NOS: 1-53 and 78-164; (d) reverse sequences of the sequences recited in SEQ ID NOS: 1-53 and 78-164; and (e) sequences having either 40%, 60%, 75% or 90% identity, as defined herein, to a sequence of (a) – (d) or a specified region of a sequence of (a) – (d).

10 In a further aspect, isolated polypeptides encoded by the polynucleotides of the present invention are provided. In one embodiment, such polypeptides comprise an amino acid sequence encoded by polynucleotides of the present invention, including polynucleotides comprising a sequence set out in the group consisting of SEQ ID NOS: 1-53 and 78-164, as well as polypeptides comprising an amino acid sequence recited in SEQ
 15 ID NOS: 165- 304.

In another aspect, the invention provides genetic constructs comprising a polynucleotide of the present invention, either alone, in combination with one or more additional polynucleotides of the present invention, or in combination with one or more known polynucleotides, together with transgenic cells comprising such constructs.

20 In a related aspect, the present invention provides genetic constructs comprising, in the 5'-3' direction, a gene promoter sequence; an open reading frame coding for at least a functional portion of an enzyme encoded by an inventive polynucleotide or a variant thereof; and a gene termination sequence. The open reading frame may be oriented in either a sense or antisense direction. Genetic constructs comprising a non-coding region

of a gene coding for an enzyme encoded by the above polynucleotide or a nucleotide sequence complementary to a non-coding region, together with a gene promoter sequence and a gene termination sequence, are also provided. Genetic constructs comprising, in the 5' – 3' direction, a promoter sequence; a polynucleotide sequence comprising at least one
5 of the following: (1) a polynucleotide comprising a polynucleotide of the present invention; or (2) a polynucleotide comprising a polynucleotide of the present invention and including a non-coding region of a gene coding for a polypeptide having activity in an isoprenoid biosynthetic pathway, are also contemplated. The genetic construct may further include a marker for the identification of transformed cells.

10 In a further aspect, transgenic host cells, such as transgenic plant cells, comprising the genetic constructs of the present invention are provided, together with plants comprising such transgenic cells, and fruits, seeds, and progeny of such plants. Other useful host cells include bacterial cells, insect cells, yeast cells and mammalian cells.

In yet another aspect, methods for modulating the isoprenoid content, composition,
15 and metabolism of an organism are provided, such methods including stably incorporating into the genome of the organism a genetic construct of the present invention. In a preferred embodiment, the target organism is a plant and the plant is a woody plant, preferably selected from the group consisting of eucalyptus, pine, acacia, poplar, sweetgum, teak and mahogany species, more preferably from the group consisting of pine
20 and eucalyptus species, and most preferably from the group consisting of *Eucalyptus grandis* and *Pinus radiata*. In a related aspect, a method for producing an organism having modified isoprenoid content is provided, the method comprising transforming a host cell with a genetic construct of the present invention to provide a transgenic cell and cultivating the transgenic cell under conditions conducive to growth and regeneration.

25 In yet a further aspect, the present invention provides methods for modifying the activity of a polypeptide in a target organism such as a plant, comprising stably incorporating into the genome of the organism a genetic construct of the present invention. In a preferred embodiment, the target organism is a plant, and the plant is a woody plant, preferably selected from the group consisting of eucalyptus, pine, acacia, poplar,
30 sweetgum, teak and mahogany species, more preferably from the group consisting of pine and eucalyptus species, and most preferably from the group consisting of *Eucalyptus grandis* and *Pinus radiata*.

In yet a further aspect, the present invention provides methods for modulating one or more of the content, the composition and the metabolism of an isoprenoid compound in an organism by administering an isolated polypeptide of the present invention to the organism. In applications in which the organism is a plant, administration of the polypeptide may be topical, such as by spraying or similar topical application. In applications in which the organism is mammalian, administration of the polypeptide may be systemic, such as by injection, intradermal delivery, oral delivery, delivery via nasal passageways or airways, or the like.

The above-mentioned and additional features of the present invention and the manner of obtaining them will become apparent, and the invention will be best understood by reference to the following more detailed description.

Description of Drawings

Fig. 1 shows a schematic diagram illustrating basic biosynthetic pathways of isoprene compounds.

Fig. 2 illustrates genomic DNA samples from tobacco plants created in a tagging experiment using a unique sequence identifier from *Pinus* (left panel) and a unique sequence identifier from *Eucalyptus* (right panel). In both panels, Lanes A and B contain DNA samples from empty-vector transformed control plants and Lanes C-E contain DNA samples from plants transformed with a unique sequence identifier.

Fig. 3 illustrates detection of a *Pinus* unique sequence identifier in transformed tobacco plants. Lanes A and B show the hybridization of a probe from SEQ ID NO: 76 to the genomic DNA of tobacco plants which lack the *Pinus* unique sequence identifier (empty-vector transformed control plants). Lanes C-E show the hybridization of the probe to the genomic DNA of tobacco plants containing one to three copies of the *Pinus* unique sequence identifier.

Fig. 4 illustrates detection of a *Eucalyptus* unique sequence identifier in transformed tobacco plants. Lanes A and B show the hybridization of a probe from SEQ ID NO: 77 to the genomic DNA of tobacco plants which lack the *Eucalyptus* unique sequence identifier (empty-vector transformed control plants). Lanes C-E show the hybridization of the probe to the genomic DNA of tobacco plants containing one to two copies of the *Eucalyptus* unique sequence identifier.

Detailed Description

As described above, isoprenoids are important components in a variety of eukaryotic functions. Modification of isoprenoid content, composition, and metabolism in the earlier parts of the pathway, especially the steps up to the formation of isopentenyl-diphosphate (IPP), geranyl-diphosphate (GPP), farnesyl-diphosphate (FPP) and squalene, may have a profound influence on the synthesis of the isoprenoid compounds deriving from these two precursors. Blocking one or more of the downstream steps branching from isopentenyl-diphosphate and squalene may also have a substantial effect on the pool of isopentenyl-diphosphate and squalene available for synthesis of terpenes or steroids. Hence, modifications in the synthesis, content, composition, and metabolism of any single enzyme in the isoprenoid biosynthetic pathway, and particularly in the early part of the pathway (IPP => GPP => FPP => squalene) of the isoprenoid synthesis, may affect the content, composition and metabolism of terpenoid and steroid compounds.

Using the methods and materials of the present invention, the isoprenoid content of a plant may be modified by incorporating sense or antisense copies of polynucleotides encoding polypeptides involved in the synthesis of isoprenoids into the genome of a target organism. In addition, the number of copies and combination of polynucleotides encoding for different enzymes in the biosynthetic pathway of isoprenoids may be manipulated to modify the relative amounts of isoprenoids synthesized, thereby producing biological materials having an altered composition and/or altered isoprenoid metabolism. Similarly, the alteration of isoprenoid composition, for direct application in a target organism, or for production of polypeptides for separate use, is advantageous for a variety of applications, as evidenced by the references cited above and incorporated herein by reference.

According to one embodiment, the present invention provides isolated polynucleotides encoding, or partially encoding, polypeptides having similarity to polypeptides known to be involved in isoprenoid synthesis and modification. The polynucleotides of the present invention were isolated from eucalyptus and pine species, but may alternatively be isolated from other plant sources and may be synthesized using conventional synthesis techniques. Specifically, isolated polynucleotides of the present invention comprise: the polynucleotides identified as SEQ ID NOS: 1-53 and 78-164; complements of the sequences identified as SEQ ID NOS: 1-53 and 78-164; reverse sequences of the sequences identified as SEQ ID NOS: 1-53 and 78-164; reverse complements of the sequences identified as SEQ ID NOS: 1-53 and 78-164; at least a

specified number of contiguous residues (*x*-mers) of any of the above-mentioned polynucleotides; polynucleotides complementary to any of the above polynucleotides; anti-sense sequences corresponding to any of the above polynucleotides; and variants of any of the above polynucleotides, as that term is described in this specification.

5 The isolated polynucleotides recited in SEQ ID NOS: 1-53 and 78-164 encode, or partially encode, polypeptides demonstrating sequence similarity to polypeptides known to be involved in an isoprenoid biosynthetic pathway, as indicated in Table 1 above. More specifically, the isolated polynucleotides listed in the first column of Table 1 encode, or partially encode the polypeptides listed in alignment in the second column of Table 1,
10 above. Predicted amino acid sequences corresponding to the polynucleotides set out in SEQ ID NOS: 1-53, 78-164, based on information available at the time of filing this application, are provided in SEQ ID NOS: 165-304, as indicated in Table 1.

 The term "polynucleotide(s)," as used herein, means a single or double-stranded polymer of deoxyribonucleotide or ribonucleotide bases and includes DNA and
15 corresponding RNA molecules, including HnRNA and mRNA molecules, both sense and anti-sense strands, and comprehends cDNA, genomic DNA and recombinant DNA, as well as wholly or partially synthesized polynucleotides. An HnRNA molecule contains introns and corresponds to a DNA molecule in a generally one-to-one manner. An mRNA molecule corresponds to an HnRNA and DNA molecule from which the introns have been
20 excised. A polynucleotide may consist of an entire gene, or any portion thereof. A gene is a polypeptide that codes for a functional polypeptide or RNA molecule. Operable anti-sense polynucleotides may comprise a fragment of the corresponding polynucleotide, and the definition of "polynucleotide" therefore includes all such operable anti-sense fragments. Anti-sense polynucleotides and techniques involving anti-sense
25 polynucleotides are well known in the art and are described, for example, in Robinson-Benion *et al.*, *Methods in Enzymol.* 254(23):363-375, 1995; and Kawasaki *et al.*, *Artific. Organs* 20(8):836-848, 1996. Polynucleotides of the present invention also encompass polynucleotide sequences that differ from the disclosed sequences but which, as a result of the degeneracy of genetic code, encode a polypeptide which is the same as that encoded
30 by a polynucleotide of the present invention.

 The definitions of the terms "complement," "reverse complement," and "reverse sequence," as used herein, are best illustrated by the following examples. For the

sequence 5' AGGACC 3', the complement, reverse complement, and reverse sequences are as follows:

	complement	3' TCCTGG 5'
	reverse complement	3' GGTCCT 5'
5	reverse sequence	5' CCAGGA 3'

Identification of genomic DNA and heterologous species DNAs can be accomplished by standard DNA/DNA hybridization techniques, under appropriately stringent conditions, using all or part of a cDNA sequence as a probe to screen an appropriate library. Alternatively, PCR techniques using oligonucleotide primers that are designed based on known genomic DNA, cDNA and protein sequences can be used to amplify and identify genomic and cDNA sequences. Synthetic DNAs corresponding to the identified sequences and variants may be produced by conventional synthesis methods. All of the polynucleotides described herein are isolated and purified, as those terms are commonly used in the art.

In another aspect, the present invention provides isolated polypeptides encoded, or partially encoded, by the above polynucleotides. As used herein, the term "polypeptide" encompasses amino acid chains of any length, including full length proteins, wherein the amino acid residues are linked by covalent peptide bonds. The term "polypeptide encoded by a polynucleotide" as used herein, includes polypeptides encoded by a polynucleotide which comprises an isolated polypeptide or variant provided herein. In one embodiment, polypeptides of the present invention comprise an amino acid sequence selected from the group consisting of sequences provided in SEQ ID NOS: 165-304, as well as variants of such sequences. According to another embodiment, polypeptides of the present invention comprise at least a specified number of contiguous residues (x-mers) of any of the sequences provided in SEQ ID NOS: 165-304.

Polypeptides of the present invention may be produced recombinantly by inserting a polynucleotide that encodes the polypeptide into an expression vector and expressing the polypeptide in an appropriate host. Any of a variety of expression vectors known to those of ordinary skill in the art may be employed. Expression may be achieved in any appropriate host cell that has been transformed or transfected with an expression vector containing a polypeptide encoding a recombinant polypeptide. Suitable host cells include prokaryotes, yeast and higher eukaryotic cells. Preferably, the host cells employed are *Escherichia coli*, insect, yeast or a mammalian cell line such as COS or CHO. The

polynucleotide(s) expressed in this manner may encode naturally occurring polypeptides, portions of naturally occurring polypeptides, or other variants thereof.

In a related aspect, polypeptides are provided that comprise at least a functional portion of a polypeptide having an amino acid sequence selected from the group consisting
5 of sequences provided in SEQ ID NOS: 165-304, and variants thereof. As used herein, a “functional portion” of a polypeptide is that portion which contains the active site essential for affecting the function of the polypeptide, for example, the portion of the molecule that is capable of binding one or more reactants. The active site may be made up of separate portions present on one or more polypeptide chains and will generally exhibit high binding
10 affinity.

Functional portions of a polypeptide may be identified by first preparing fragments of the polypeptide by either chemical or enzymatic digestion of the polypeptide, or by mutation analysis of the polynucleotide that encodes the polypeptide and subsequent expression of the resulting mutant polypeptides. The polypeptide fragments or mutant
15 polypeptides are then tested to determine which portions retain biological activity, using, for example, the representative assays provided below.

A functional portion comprising an active site may be made up of separate portions present on one or more polypeptide chains and generally exhibits high substrate specificity. The term “polypeptide encoded by a polynucleotide” as used herein, includes
20 polypeptides encoded by a polynucleotide comprising a partial isolated polynucleotide of the present invention.

Portions and other variants of the inventive polypeptides may also be generated by synthetic or recombinant means. Synthetic polypeptides having fewer than about 100 amino acids, and generally fewer than about 50 amino acids, may be generated using
25 techniques that are well known to those of ordinary skill in the art. For example, such polypeptides may be synthesized using any of the commercially available solid-phase techniques, such as the Merrifield solid-phase synthesis method, where amino acids are sequentially added to a growing amino acid chain. See Merrifield, *J. Am. Chem. Soc.* 85:2149-2154, 1963. Equipment for automated synthesis of polypeptides is commercially
30 available from suppliers such as Perkin Elmer/Applied Biosystems, Inc. (Foster City, CA), and may be operated according to the manufacturer's instructions. Variants of a native polypeptide may be prepared using standard mutagenesis techniques, such as oligonucleotide-directed site-specific mutagenesis (Kunkel T, *Proc. Natl. Acad. Sci. USA*

82: 488-492, 1985). Sections of DNA sequences may also be removed using standard techniques to permit preparation of truncated polypeptides.

In general, the polypeptides disclosed herein are prepared in an isolated, substantially pure form. Preferably, the polypeptides are at least about 80% pure; more preferably at least about 90% pure; and most preferably, at least about 99% pure. In certain preferred embodiments, described in detail below, the isolated polypeptides are incorporated into pharmaceutical compositions or vaccines for use in the treatment of skin disorders.

As used herein, the term "variant" comprehends polynucleotide or polypeptide sequences different from the specifically identified sequences, wherein one or more nucleotides or amino acid residues is deleted, substituted, or added. Variants may be naturally occurring allelic variants, or non-naturally occurring variants. Variant polynucleotide sequences preferably exhibit at least 40%; more preferably at least 60%; more preferably yet at least 75%; and most preferably at least 90% identity to a sequence of the present invention. Variant polypeptide sequences preferably exhibit at least 50%; more preferably at least 75%; more preferably yet at least 90%; and most preferably at least 95% identity to a sequence of the present invention. The percentage identity is determined by aligning the two sequences to be compared as described below, determining the number of identical residues in the aligned portion, dividing that number by the total number of residues in the inventive (queried) sequence, and multiplying the result by 100.

Polynucleotide and polypeptide sequences may be aligned, and percentage of identical residues in a specified region may be determined against another polynucleotide or polypeptide, using computer algorithms that are publicly available. Two exemplary algorithms for aligning and identifying the similarity of polynucleotide sequences are the BLASTN and FASTA algorithms. Polynucleotides may also be analyzed using the BLASTX algorithm, which compares the six-frame conceptual translation products of a nucleotide query sequence (both strands) against a protein sequence database. The percentage identity of polypeptide sequences may be examined using the BLASTP algorithm. The BLASTN, BLASTX and BLASTP programs are available on the NCBI anonymous FTP server (<ftp://ncbi.nlm.nih.gov>) under /blast/executables/. The BLASTN algorithm Version 2.0.4 [Feb-24-1998] and Version 2.0.6 [Sept-16-1998], set to the parameters described below, is preferred for use in the determination of polynucleotide variants according to the present invention. The BLASTP algorithm, set to the parameters

described below, is preferred for use in the determination of polypeptide variants according to the present invention. The use of the BLAST family of algorithms, including BLASTN, BLASTP, and BLASTX, is described at NCBI's website at URL <http://www.ncbi.nlm.nih.gov/BLAST/newblast.html> and in the publication of Altschul, *et al.*, *Nucleic Acids Res.* 25: 3389-3402, 1997.

The computer algorithm FASTA is available on the Internet at the ftp site <ftp://ftp.virginia.edu/pub/fasta/>. Version 2.0u4 [February 1996], set to the default parameters described in the documentation and distributed with the algorithm, may be also used in the determination of variants according to the present invention. The use of the FASTA algorithm is described in Pearson and Lipman, *Proc. Natl. Acad. Sci. USA* 85:2444-2448, 1988; and Pearson WR, *Methods in Enzymol.* 183: 63-98, 1990.

The following running parameters are preferred for determination of alignments and similarities using BLASTN that contribute to the E values and percentage identity for polynucleotide sequences: Unix running command: `blastall -p blastn -d embldb -e 10 -G0 -E0 -r 1 -v 30 -b 30 -i queryseq -o results`; the parameters are: -p Program Name [String]; -d Database [String]; -e Expectation value (E) [Real]; -G Cost to open a gap (zero invokes default behavior) [Integer]; -E Cost to extend a gap (zero invokes default behavior) [Integer]; -r Reward for a nucleotide match (BLASTN only) [Integer]; -v Number of one-line descriptions (V) [Integer]; -b Number of alignments to show (B) [Integer]; -i Query File [File In]; and -o BLAST report Output File [File Out] Optional.

The following running parameters are preferred for determination of alignments and similarities using BLASTP that contribute to the E values and percentage identity of polypeptide sequences: `blastall -p blastp -d swissprot -e 10 -G 0 -E 0 -v 30 -b 30 -i queryseq -o results`; the parameters are: -p Program Name [String]; -d Database [String]; -e Expectation value (E) [Real]; -G Cost to open a gap (zero invokes default behavior) [Integer]; -E Cost to extend a gap (zero invokes default behavior) [Integer]; -v Number of one-line descriptions (v) [Integer]; -b Number of alignments to show (b) [Integer]; -i Query File [File In]; -o BLAST report Output File [File Out] Optional. The "hits" to one or more database sequences by a queried sequence produced by BLASTN, FASTA, BLASTP or a similar algorithm, align and identify similar portions of sequences. The hits are arranged in order of the degree of similarity and the length of sequence overlap. Hits to a database sequence generally represent an overlap over only a fraction of the sequence length of the queried sequence.

The BLASTN, FASTA, and BLASTP algorithms also produce "Expect" values for alignments. The Expect value (E) indicates the number of hits one can "expect" to see over a certain number of contiguous sequences by chance when searching a database of a certain size. The Expect value is used as a significance threshold for determining whether
5 the hit to a database, such as the preferred EMBL database, indicates true similarity. For example, an E value of 0.1 assigned to a polynucleotide hit is interpreted as meaning that in a database of the size of the EMBL database, one might expect to see 0.1 matches over the aligned portion of the sequence with a similar score simply by chance. By this criterion, the aligned and matched portions of the polynucleotide sequences then have a
10 probability of 90% of being the same. For sequences having an E value of 0.01 or less over aligned and matched portions, the probability of finding a match by chance in the EMBL database is 1% or less using the BLASTN or FASTA algorithm.

According to one embodiment, "variant" polynucleotides and polypeptides, with reference to each of the polynucleotides and polypeptides of the present invention,
15 preferably comprise sequences producing an E value of 0.01 or less when compared to the polynucleotide or polypeptide of the present invention. That is, a variant polynucleotide or polypeptide is any sequence that has at least a 99% probability of being the same as the polynucleotide or polypeptide of the present invention, measured as having an E value of 0.01 or less using the BLASTN, FASTA, or BLASTP algorithms set at parameters
20 described above. According to a preferred embodiment, a variant polynucleotide is a sequence having the same number or fewer nucleic acids than a polynucleotide of the present invention that has at least a 99% probability of being the same as the polynucleotide of the present invention, measured as having an E value of 0.01 or less using the BLASTN or FASTA algorithms set at parameters described above. Similarly,
25 according to a preferred embodiment, a variant polypeptide is a sequence having the same number or fewer amino acids than a polypeptide of the present invention that has at least a 99% probability of being the same as a polypeptide of the present invention, measured as having an E value of 0.01 or less using the BLASTP algorithm set at the parameters described above.

30 Alternatively, variant polynucleotides or polypeptides of the present invention comprise a sequence exhibiting at least 40%; more preferably at least 60%; more preferably yet at least 75%; and most preferably at least 90% identity to a polynucleotide or polypeptide of the present invention, determined as described below. The percentage

identity is determined by aligning sequences using one of the BLASTN, FASTA, or BLASTP algorithms, set at the running parameters described above, and identifying the number of identical nucleic or amino acids over the aligned portions; dividing the number of identical nucleic or amino acids by the total number of nucleic or amino acids of the polynucleotide or polypeptide of the present invention; and then multiplying by 100 to determine the percentage identity. For example, a polynucleotide of the present invention having 220 nucleic acids has a hit to a polynucleotide sequence in the EMBL database having 520 nucleic acids over a stretch of 23 nucleotides in the alignment produced by the BLASTN algorithm using the parameters described above. The 23 nucleotide hit includes 21 identical nucleotides, one gap and one different nucleotide. The percentage identity of the polynucleotide of the present invention to the hit in the EMBL library is thus 21/220 times 100, or 9.5%. The polynucleotide sequence in the EMBL database is thus not a variant of a polynucleotide of the present invention.

Alternatively, variant polynucleotides of the present invention hybridize to the polynucleotide sequences recited in SEQ ID NOS: 1-53 and 78-164, or complements, reverse sequences, or reverse complements of those sequences under stringent conditions. As used herein, "stringent conditions" refers to prewashing in a solution of 6X SSC, 0.2% SDS; hybridizing at 65°C, 6X SSC, 0.2% SDS overnight; followed by two washes of 30 minutes each in 1X SSC, 0.1% SDS at 65°C and two washes of 30 minutes each in 0.2X SSC, 0.1% SDS at 65°C.

The present invention also encompasses polynucleotides that differ from the disclosed sequences but that, as a consequence of the discrepancy of the genetic code, encode a polypeptide having similar enzymatic activity as a polypeptide encoded by a polynucleotide of the present invention. Thus, polynucleotides comprising sequences that differ from the polynucleotide sequences recited in SEQ ID NOS: 1-53 and 78-164, or complements, reverse sequences, or reverse complements of those sequences as a result of conservative substitutions are contemplated by and encompassed within the present invention. Additionally, polynucleotides comprising sequences that differ from the polynucleotide sequences recited in SEQ ID NOS: 1-53 and 78-164, or complements, reverse complements, or reverse sequences as a result of deletions and/or insertions totaling less than 10% of the total sequence length are also contemplated by and encompassed within the present invention. Similarly, polypeptides comprising sequences that differ from the polypeptide sequences recited in SEQ ID NOS: 165-304 as a result of

amino acid substitutions, insertions, and/or deletions totaling less than 10% of the total sequence length are contemplated by an encompassed within the present invention, provided the variant polypeptide has activity in an isoprenoid biosynthetic pathway.

The polynucleotides of the present invention may be isolated from various libraries, or may be synthesized using techniques that are well known in the art. The polynucleotides may be synthesized, for example, using automated oligonucleotide synthesizers (e.g., Beckman Oligo 1000M DNA Synthesizer) to obtain polynucleotide segments of up to 50 or more nucleic acids. A plurality of such polynucleotide segments may then be ligated using standard DNA manipulation techniques that are well known in the art of molecular biology. One conventional and exemplary polynucleotide synthesis technique involves synthesis of a single stranded polynucleotide segment having, for example, 80 nucleic acids, and hybridizing that segment to a synthesized complementary 85 nucleic acid segment to produce a 5 nucleotide overhang. The next segment may then be synthesized in a similar fashion, with a 5 nucleotide overhang on the opposite strand. The "sticky" ends ensure proper ligation when the two portions are hybridized. In this way, a complete polynucleotide of the present invention may be synthesized entirely *in vitro*.

Some of the polynucleotides identified as SEQ ID NOS: 1-53 and 78-164 are referred to as "partial" sequences, in that they do not represent the full coding portion of a gene encoding a naturally occurring polypeptide. The partial polynucleotide sequences disclosed herein may be employed to obtain the corresponding full length genes for various species and organisms by, for example, screening DNA expression libraries using hybridization probes based on the polynucleotides of the present invention, or using PCR amplification with primers based upon the polynucleotides of the present invention. In this way one can, using methods well known in the art, extend a polynucleotide of the present invention upstream and downstream of the corresponding mRNA, as well as identify the corresponding genomic DNA, including the promoter and enhancer regions, of the complete gene. The present invention thus comprehends isolated polynucleotides comprising a sequence identified in SEQ ID NOS: 1-53 and 78-164, or a variant of one of the specified sequences, that encode a functional polypeptide, including full length genes. Such extended polynucleotides may have a length of from about 50 to about 4,000 nucleic acids or base pairs, and preferably have a length of less than about 4,000 nucleic acids or base pairs, more preferably yet a length of less than about 3,000 nucleic acids or base

pairs, more preferably yet a length of less than about 2,000 nucleic acids or base pairs. Under some circumstances, extended polynucleotides of the present invention may have a length of less than about 1,800 nucleic acids or base pairs, preferably less than about 1,600 nucleic acids or base pairs, more preferably less than about 1,400 nucleic acids or base
5 pairs, more preferably yet less than about 1,200 nucleic acids or base pairs, and most preferably less than about 1,000 nucleic acids or base pairs.

Polynucleotides of the present invention also comprehend polynucleotides comprising at least a specified number of contiguous residues (x -mers) of any of the polynucleotides identified as SEQ ID NOS: 1-53 and 78-164, complements, reverse
10 sequences, and reverse complements of such sequences, and their variants. Similarly, polypeptides of the present invention comprehend polypeptides comprising at least a specified number of contiguous residues (x -mers) of any of the polypeptides identified as SEQ ID NOS: 165-304, and their variants. As used herein, the term " x -mer," with reference to a specific value of " x ," refers to a sequence comprising at least a specified
15 number (" x ") of contiguous residues of any of the polynucleotides identified as SEQ ID NOS: 1-53 and 78-164, or the polypeptides identified as SEQ ID NOS: 165-304. According to preferred embodiments, the value of x is preferably at least 20; more preferably, at least 40; more preferably yet, at least 60; and most preferably, at least 80. Thus, polynucleotides and polypeptides of the present invention comprise a 20-mer, a 40-
20 mer, a 60-mer, an 80-mer, a 100-mer, a 120-mer, a 150-mer, a 180-mer, a 220-mer, a 250-mer, or a 300-mer, 400-mer, 500-mer or 600-mer of a polynucleotide or polypeptide identified as SEQ ID NOS: 1-53, and 78-304, and variants thereof.

Polynucleotide probes and primers complementary to and/or corresponding to SEQ ID NOS: 1-53 and 78-164, and variants of those sequences, are also comprehended by the
25 present invention. Such oligonucleotide probes and primers are substantially complementary to the polynucleotide of interest. As used herein, the term "oligonucleotide" refers to a relatively short segment of a polynucleotide sequence, generally comprising between 6 and 60 nucleotides, and comprehends both probes for use in hybridization assays and primers for use in the amplification of DNA by polymerase
30 chain reaction.

An oligonucleotide probe or primer is described as "corresponding to" a polynucleotide of the present invention, including one of the sequences set out as SEQ ID NOS: 1-53 and 78-164, or a variant, if the oligonucleotide probe or primer, or its

complement, is contained within one of the sequences set out as SEQ ID NOS: 1-53 and 78-164, or a variant of one of the specified sequences.

Two single stranded sequences are said to be substantially complementary when the nucleotides of one strand, optimally aligned and compared, with the appropriate
5 nucleotide insertions and/or deletions, pair with at least 80%, preferably at least 90% to 95%, and more preferably at least 98% to 100%, of the nucleotides of the other strand. Alternatively, substantial complementarity exists when a first DNA strand selectively hybridizes to a second DNA strand under stringent hybridization conditions. Stringent hybridization conditions for determining complementarity include salt conditions of less
10 than about 1 M, more usually less than about 500 mM and preferably less than about 200 mM. Hybridization temperatures may be as low as 5°C, but are generally greater than about 22°C, more preferably greater than about 30°C and most preferably greater than about 37°C. Longer DNA fragments may require higher hybridization temperatures for specific hybridization. Since the stringency of hybridization may be affected by other
15 factors such as probe composition, presence of organic solvents and extent of base mismatching, the combination of parameters is more important than the absolute measure of any one alone. The DNA from plants or samples or products containing plant material can be either genomic DNA or DNA derived by preparing cDNA from the RNA present in the sample.

20 In addition to DNA-DNA hybridization, DNA-RNA or RNA-RNA hybridization assays are also possible. In the first case, the mRNA from expressed genes would then be detected instead of genomic DNA or cDNA derived from mRNA of the sample. In the second case, RNA probes could be used. In addition, artificial analogs of DNA hybridizing specifically to target sequences could also be used.

25 In specific embodiments, the oligonucleotide probes and/or primers comprise at least about 6 contiguous residues, more preferably at least about 10 contiguous residues, and most preferably at least about 20 contiguous residues complementary to a polynucleotide sequence of the present invention. Probes and primers of the present invention may be from about 8 to 100 base pairs in length or, preferably, from about 10 to
30 50 base pairs in length or, more preferably, from about 15 to 40 base pairs in length. The probes can be easily selected using procedures well known in the art, taking into account DNA-DNA hybridization stringencies, annealing and melting temperatures, potential for formation of loops and other factors, which are well known in the art. Tools and software

suitable for designing probes, and especially suitable for designing PCR primers, are available on the Internet, for example, at URL <http://www.horizonpress.com/pcr/>. Preferred techniques for designing PCR primers are also disclosed in Dieffenbach CW and Dyksler GS, *PCR primer: a laboratory manual*. CSHL Press: Cold Spring Harbor, NY, 5 1995.

A plurality of oligonucleotide probes or primers corresponding to a polynucleotide of the present invention may be provided in a kit form. Such kits generally comprise multiple DNA or oligonucleotide probes, each probe being specific for a polynucleotide sequence. Kits of the present invention may comprise one or more probes or primers 10 corresponding to a polynucleotide of the present invention, including a polynucleotide sequence identified in SEQ ID NOS: 1-53 and 78-164.

In one embodiment useful for high-throughput assays, the oligonucleotide probe kits of the present invention comprise multiple probes in an array format, wherein each probe is immobilized in a predefined, spatially addressable location on the surface of a 15 solid substrate. Array formats which may be usefully employed in the present invention are disclosed, for example, in U.S. Patent Nos. 5,412,087 and 5,545,531; and PCT Publication No. WO 95/00530, the disclosures of which are hereby incorporated by reference.

Probes, preferably in the form of an array, may be employed to screen for 20 differences in organisms or samples or products containing genetic material using high throughput screening techniques that are well known in the art. The significance of using probes in high-throughput screening systems is apparent for applications such as plant breeding and quality control operations in which there is a need to identify large numbers of seed lots and plant seedlings, to examine samples or products for unwanted plant 25 materials, to identify plants or samples or products containing plant material for quarantine purposes, etc., or to ascertain the true origin of plants or samples or products containing plant material. Screening for the presence or absence of polynucleotides of the present invention used as identifiers for tagging plants is valuable for later detecting the amount of gene flow in plant breeding, introgression of genes via dispersed pollen, etc.

30 In this manner, oligonucleotide probe kits of the present invention may be employed to examine the presence/absence (or relative amounts in case of mixtures) of polynucleotides in different samples or products containing different materials rapidly and in a cost-effective manner. Examples of plant species that may be examined using the

present invention, include forestry species, such as pine and eucalyptus species, other tree species, and agricultural and horticultural plants.

Another aspect of the present invention involves collections of a plurality of polynucleotides of the present invention. A collection of a plurality of the polynucleotides
5 of the present invention, particularly the polynucleotides identified as SEQ ID NOS: 1-53 and 78-164, and variants thereof, may be recorded and/or stored on a storage medium and subsequently accessed for purposes of analysis, comparison, etc. Suitable storage media include magnetic media such as magnetic diskettes, magnetic tapes, CD-ROM storage media, optical storage media, and the like. Suitable storage media and methods for
10 recording and storing information, as well as accessing information such as polynucleotide sequences recorded on such media, are well known in the art. The polynucleotide information stored on the storage medium is preferably computer-readable and may be used for analysis and comparison of the polynucleotide information.

According to one embodiment, the storage medium includes a collection of at least
15 4, preferably at least 10, more preferably at least 15, and most preferably at least 20 of the polynucleotides of the present invention, preferably the polynucleotides identified as SEQ ID NOS: 1-53 and 78-164, and variants of those polynucleotides.

For applications where modulation of a polypeptide involved with isoprenoid biosynthesis and/or isoprenoid metabolism is desired, an open reading frame may be
20 inserted into a genetic construct in a sense or antisense orientation, such that transformation of a target plant with the genetic construct produces a change in the expression level of the polypeptide compared to the expression in a wild-type organism. Transformation with a genetic construct comprising an open reading frame in a sense orientation will generally result in modulation of expression of the selected gene, while
25 transformation with a genetic construct comprising an open reading frame in an antisense orientation generally produces reduced expression of the selected gene. A population of plants transformed with a genetic construct comprising an open reading frame of the present invention in either a sense or antisense orientation may be screened for increased or reduced expression of the gene in question using techniques well known to those of
30 skill in the art, and plants having the desired phenotypes may thus be isolated.

Alternatively, expression of a gene involved in the biosynthesis of isoprenoids may be inhibited by inserting a portion of an open reading frame of the present invention, in either sense or antisense orientation, in the genetic construct. Such portions need not be

full-length but preferably comprise at least 25, and more preferably, at least 50 residues of polynucleotide of the present invention. A much longer portion, or even the full length polynucleotide corresponding to the complete open reading frame, may be employed. The portion of the open reading frame does not need to be precisely the same as the
5 endogenous sequence, provided that there is sufficient sequence similarity to achieve inhibition of the target gene. Thus a sequence derived from one species may be used to inhibit expression of a gene in a different species.

According to another embodiment, the genetic constructs of the present invention comprise a polynucleotide including a non-coding region of a gene coding for a polypeptide encoded by a polynucleotide of the present invention, or a polynucleotide
10 complementary to such a non-coding region. Examples of non-coding regions which may be usefully employed in such constructs include introns and 5'-non-coding leader sequences. Transformation of a target plant with such a genetic construct may lead to a reduction in the amount of an isoprenoid compound synthesized by the plant by the
15 process of cosuppression, in a manner similar to that discussed, for example, by Napoli *et al.*, *Plant Cell* 2:279-290, 1990 and de Carvalho Niebel *et al.*, *Plant Cell* 7:347-358, 1995.

Alternatively, regulation may be achieved by inserting appropriate sequences or subsequences (e.g. DNA or RNA) in ribozyme constructs (McIntyre CL and Manners JM, *Transgenic Res.* 5(4):257-262, 1996). Ribozymes are synthetic RNA molecules that
20 comprise a hybridizing region complementary to two regions, each of which comprises at least 5 contiguous nucleotides in a mRNA molecule encoded by one of the inventive polynucleotides. Ribozymes possess highly specific endonuclease activity, which autocatalytically cleaves the mRNA.

The genetic constructs of the present invention further comprise a gene promoter
25 sequence and a gene termination sequence, operably linked to the polynucleotide to be transcribed, which control expression of the polypeptide. The gene promoter sequence is generally positioned at the 5' end of the polynucleotide to be transcribed, and is employed to initiate transcription of the polynucleotide. Gene promoter sequences are generally found in the 5' non-coding region of a gene but they may exist downstream of the open
30 reading frame or in introns (Luehrsen KR, *Mol. Gen. Genet.* 225:81-93, 1991); or in the coding region, as for example in a plant defence gene (Douglas *et al.*, *EMBO J.* 10:1767-1775, 1991). When the construct includes an open reading frame in a sense orientation, the gene promoter sequence also initiates translation of the open reading frame. For

genetic constructs comprising either an open reading frame in an antisense orientation or a non-coding region, the gene promoter sequence consists only of a transcription initiation site having a RNA polymerase binding site.

Numerous gene promoter sequences that may be usefully employed in the genetic constructs of the present invention are well known in the art. The gene promoter sequence, and also the gene termination sequence, may be endogenous to the target plant host or may be exogenous, provided the promoter is functional in the target host. For example, the promoter and termination sequences may be from other plant species, plant viruses, bacterial plasmids and the like. Preferably, gene promoter and termination sequences are common to those of the polynucleotide being introduced.

Factors influencing the choice of promoter include the desired tissue specificity of the construct, and the timing of transcription and translation. For example, constitutive promoters, such as the 35S Cauliflower Mosaic Virus (CaMV 35S) promoter with or without enhancers, such as the Kozak sequence or the Omega enhancer, and *Agrobacterium tumefaciens* nopaline synthase terminator, may be usefully employed in the present invention. Use of a tissue specific promoter will result in production of the desired sense or antisense RNA only in the tissue of interest. With genetic constructs employing inducible gene promoter sequences, the rate of RNA polymerase binding and initiation can be modulated by external stimuli, such as light, heat, anaerobic stress, alteration in nutrient conditions and the like. Temporally regulated promoters can be employed to effect modulation of the rate of RNA polymerase binding and initiation at a specific time during development of a transformed cell. Preferably, the original promoters from the enzyme gene in question, or promoters from a specific tissue-targeted gene in the organism to be transformed, such as eucalyptus or pine are used. Other examples of gene promoters which may be usefully employed in the present invention include mannopine synthase (mas), octopine synthase (ocs) and those reviewed by Chua *et al.*, *Science* 244:174-181, 1989.

The gene termination sequence, which is located 3' to the polynucleotide to be transcribed, may come from the same gene as the gene promoter sequence or may be from a different gene. Many gene termination sequences known in the art may be usefully employed in the present invention, such as the 3' end of the *Agrobacterium tumefaciens* nopaline synthase gene. However, preferred gene terminator sequences are those from the original enzyme gene or from the target species to be transformed.

The genetic constructs of the present invention may also contain a selection marker that is effective in target cells, such as plant cells, to allow for the detection of transformed cells containing the inventive construct. Such markers, which are well known in the art, typically confer resistance to one or more toxins. One example of such a marker is the NPTII gene whose expression results in resistance to kanamycin or hygromycin, antibiotics which are usually toxic to plant cells at a moderate concentration (Rogers et al. in Weissbach A and Weissbach H, eds., *Methods for Plant Molecular Biology*, Academic Press Inc.: San Diego, CA, 1988). Transformed cells can thus be identified by their ability to grow in media containing the antibiotic in question. Alternatively, the presence of the desired construct in transformed cells can be determined by means of other techniques well known in the art, such as Southern and Western blots. A transcription initiation site may additionally included in the genetic construct when the sequence to be transcribed lacks such a site.

Techniques for operatively linking the components of the genetic constructs of the present invention are well known in the art and include the use of synthetic linkers containing one or more restriction endonuclease sites as described, for example, by Sambrook et al., *Molecular cloning: a laboratory manual*, CSHL Press: Cold Spring Harbor, NY, 1989. The DNA construct of the present invention may be linked to a vector having at least one replication system, for example *E. coli*, whereby after each manipulation, the resulting construct can be cloned and sequenced and the correctness of the manipulation determined.

The genetic constructs of the present invention may be used to transform a variety of target organisms such as plants, both monocotyledonous (e.g., grasses, corn, grains, oat, wheat and barley); dicotyledonous (e.g., *Arabidopsis*, tobacco, legumes, alfalfa, oaks, eucalyptus, maple); gymnosperms (e.g., Scots pine (Aronen, *Finnish Forest Res. Papers*, Vol. 595, 1996); white spruce (Ellis et al., *Biotechnology* 11: 84-89, 1993); and larch (Huang et al., *In Vitro Cell* 27:201-207, 1991). In a preferred embodiment, the inventive DNA constructs are employed to transform woody plants, herein defined as a tree or shrub whose stem lives for a number of years and increases in diameter each year by the addition of woody tissue. Preferably the target plant is selected from the group consisting of eucalyptus and pine species, most preferably from the group consisting of *Eucalyptus grandis* and *Pinus radiata*. Other species which may be usefully transformed with the DNA constructs of the present invention include, but are not limited to: Pines, such as

Pinus banksiana, *Pinus brutia*, *Pinus caribaea*, *Pinus clausa*, *Pinus contorta*, *Pinus coulteri*, *Pinus echinata*, *Pinus eldarica*, *Pinus ellioti*, *Pinus jeffreyi*, *Pinus lambertiana*, *Pinus monticola*, *Pinus nigra*, *Pinus palustris*, *Pinus pinaster*, *Pinus ponderosa*, *Pinus resinosa*, *Pinus rigida*, *Pinus serotina*, *Pinus strobus*, *Pinus sylvestris*, *Pinus taeda*, *Pinus virginiana*; other gymnosperm, such as *Abies amabilis*, *Abies balsamea*, *Abies concolor*, *Abies grandis*, *Abies lasiocarpa*, *Abies magnifica*, *Abies procera*, *Chamaecyparis lawsoniana*, *Chamaecyparis nootkatensis*, *Chamaecyparis thyoides*, *Huniperus virginiana*, *Larix decidua*, *Larix laricina*, *Larix leptolepis*, *Larix occidentalis*, *Larix siberica*, *Libocedrus decurrens*, *Picea abies*, *Picea engelmanni*, *Picea glauca*, *Picea mariana*, *Picea pungens*, *Picea rubens*, *Picea sitchensis*, *Pseudotsuga menziesii*, *Sequoia gigantea*, *Sequoia sempervirens*, *Taxodium distichum*, *Tsuga canadensis*, *Tsuga heterophylla*, *Tsuga mertensiana*, *Thuja occidentalis*, *Thuja plicata*; and Eucalypts, such as *Eucalyptus alba*, *Eucalyptus bancroftii*, *Eucalyptus botyroides*, *Eucalyptus bridgesiana*, *Eucalyptus calophylla*, *Eucalyptus camaldulensis*, *Eucalyptus citriodora*, *Eucalyptus cladocalyx*, *Eucalyptus coccifera*, *Eucalyptus curtisii*, *Eucalyptus dalrympleana*, *Eucalyptus deglupta*, *Eucalyptus delagatensis*, *Eucalyptus diversicolor*, *Eucalyptus dunnii*, *Eucalyptus ficifolia*, *Eucalyptus globulus*, *Eucalyptus gomphocephala*, *Eucalyptus gunnii*, *Eucalyptus henryi*, *Eucalyptus laevopinea*, *Eucalyptus macarthurii*, *Eucalyptus macrorhyncha*, *Eucalyptus maculata*, *Eucalyptus marginata*, *Eucalyptus megacarpa*, *Eucalyptus melliodora*, *Eucalyptus nicholii*, *Eucalyptus nitens*, *Eucalyptus nova-anglica*, *Eucalyptus obliqua*, *Eucalyptus obtusiflora*, *Eucalyptus oreades*, *Eucalyptus pauciflora*, *Eucalyptus polybractea*, *Eucalyptus regnans*, *Eucalyptus resinifera*, *Eucalyptus robusta*, *Eucalyptus rudis*, *Eucalyptus saligna*, *Eucalyptus sideroxylon*, *Eucalyptus stuartiana*, *Eucalyptus tereticornis*, *Eucalyptus torelliana*, *Eucalyptus urnigera*, *Eucalyptus urophylla*, *Eucalyptus viminalis*, *Eucalyptus viridis*, *Eucalyptus wandoo*, *Eucalyptus youmanni*.

Techniques for stably incorporating genetic constructs into the genome of target plants are well known in the art and include *Agrobacterium tumefaciens* mediated introduction, electroporation, protoplast fusion, injection into reproductive organs, injection into immature embryos, high velocity projectile introduction, and the like. The choice of technique will depend upon the target plant to be transformed. For example, dicotyledonous plants and certain monocots and gymnosperms may be transformed by *Agrobacterium* Ti plasmid technology, as described, for example by Bevan, *Nucleic Acid Res.* 12:8711-8721, 1984. Targets for the introduction of the genetic constructs of the

present invention include tissues, such as leaf tissue, disseminated cells, protoplasts, seeds, embryos, meristematic regions; cotyledons, hypocotyls, and the like. The preferred method for transforming eucalyptus and pine is a biolistic method using pollen (*see, for example, Aronen, Finnish Forest Res. Papers 595:53, 1996*) or easily regenerable
5 embryonic tissues.

Once the cells are transformed, cells having the inventive genetic construct incorporated in their genome may be selected by means of a marker, such as the kanamycin resistance marker discussed above. Transgenic cells may then be cultured in an appropriate medium to regenerate whole plants, using techniques well known in the art.
10 In the case of protoplasts, the cell wall is allowed to reform under appropriate osmotic conditions. In the case of seeds or embryos, an appropriate germination or callus initiation medium is employed. For explants, an appropriate regeneration medium is used. Regeneration of plants is well established for many species. For a review of regeneration of forest trees, *see Dunstan et al., in Thorpe TA, ed., In vitro embryogenesis of plants,*
15 *Current Plant Science and Biotechnology in Agriculture, 20(12):471-540, 1995.* Specific protocols for the regeneration of spruce are discussed by Roberts., "Somatic embryogenesis of spruce," in Redenbaugh K, ed., *Synseed: applications of synthetic seed to crop improvement*, CRC Press: Ch. 23, pp. 427-449, 1993. The resulting transformed plants may be reproduced sexually or asexually, using methods well known in the art, to give
20 successive generations of transgenic plants.

As discussed above, the production of RNA in target plant cells can be controlled by choice of the promoter sequence, or by selecting the number of functional copies or the site of integration of the polynucleotides incorporated into the genome of the target plant host. A target plant may be transformed with more than one genetic constructs of the
25 present invention, thereby modulating the activity of more than one isoprenoid metabolism enzyme, affecting enzyme activity in more than one tissue, or affecting enzyme activity at more than one expression time. Similarly, a genetic construct may be assembled containing more than one open reading frame coding for an enzyme encoded by a polynucleotide of the present invention or more than one non-coding region of a gene
30 coding for such an enzyme. The polynucleotides of the present inventive may also be employed in combination with other known sequences encoding enzymes involved in the synthesis of isoprenoids.

Additionally, the polynucleotides of the present invention have particular application for use as non-disruptive tags for marking organisms, particularly plants. Genetic constructs comprising polynucleotides of the present invention may be stably introduced into an organism as heterologous, non-functional, non-disruptive tags. It is
5 then possible to identify the origin or source of the organism at a later date by determining the presence or absence of the tag(s) in a sample of material. Organisms other than plants may also be tagged with the polynucleotides of the present invention, including commercially valuable animals, fish, bacteria and yeasts.

Detection of the tag(s) may be accomplished using a variety of conventional
10 techniques, and will generally involve the use of nucleic acid probes. Sensitivity in assaying the presence of probe can be usefully increased by using branched oligonucleotides, as described by Horn *et al.*, *Nucleic Acids Res.* 25(23):4842-4849, 1997), enabling detection of as few as 50 DNA molecules in the sample.

The following examples are offered by way of illustration and not by way of
15 limitation.

Example 1

Isolation and Characterization of cDNA Clones from *Pinus radiata* and *Eucalyptus grandis*

Pinus radiata and *Eucalyptus grandis* cDNA expression libraries were constructed
20 and screened as follows. mRNA was extracted from the plant tissue using the protocol of Chang *et al.*, *Plant Molecular Biology Reporter* 11:113-116, 1993 with minor modifications. Specifically, samples were dissolved in CPC-RNAXB (100 mM Tris-Cl, pH 8.0; 25 mM EDTA; 2.0 M NaCl; 2%CTAB; 2% PVP and 0.05% Spermidine*3HCl) and extracted with chloroform:isoamyl alcohol, 24:1. mRNA was precipitated with
25 ethanol and the total RNA preparate was purified using a Poly(A) Quik mRNA Isolation Kit (Stratagene, La Jolla, CA). A cDNA expression library was constructed from the purified mRNA by reverse transcriptase synthesis followed by insertion of the resulting cDNA clones in Lambda ZAP using a ZAP Express cDNA Synthesis Kit (Stratagene), according to the manufacturer's protocol. The resulting cDNAs were packaged using a
30 Gigapack II Packaging Extract (Stratagene) employing 1 µl of sample DNA from the 5 µl ligation mix. Mass excision of the library was done using XL1-Blue MRF' cells and XL0LR cells (Stratagene) with ExAssist helper phage (Stratagene). The excised phagemids were diluted with NZY broth (Gibco BRL, Gaithersburg, MD) and plated out

onto LB-kanamycin agar plates containing X-gal and isopropylthio-beta-galactoside (IPTG).

Of the colonies plated and picked for DNA miniprep, the large majority contained an insert suitable for sequencing. Positive colonies were cultured in NZY broth with kanamycin and cDNA was purified by means of REAL DNA minipreps (Qiagen, Venlo, The Netherlands). Agarose gel at 1% was used to screen sequencing templates for chromosomal contamination. Dye terminator sequences were prepared using a Biomek 2000 robot (Beckman Coulter Inc, Fullerton CA for liquid handling and DNA amplification using a 9700 PCR machine (Perkin Elmer/Applied Biosystems, Foster City, CA) according to the manufacturer's protocol.

Polynucleotides for positive clones were obtained using a Perkin Elmer/Applied Biosystems Division Prism 377 sequencer. cDNA clones were sequenced first from the 5' end and, in some cases, also from the 3' end. For some clones, internal sequences were obtained using subcloned fragments. Subcloning was performed using standard procedures of restriction mapping and subcloning to pBluescript II SK+ vector and other standard sequencing vectors.

The determined cDNA sequences, including the polynucleotides of the present invention, were compared to and aligned with known sequences in the. Specifically, the polynucleotides identified in SEQ ID NOS. 1-53 were compared to polynucleotides in the EMBL database EMBL as of the end of August, 1998 using the BLASTN algorithm Version 2.0.4 [Feb-24-1998] set to the following running: Unix running command: blastall -p blastn -d embldb -e 10 -G 0 -E 0 -r 1 -v 30 -b 30 -i queryseq -o results. The polynucleotides identified in SEQ ID NOS: 78-164 were compared to polynucleotides in the EMBL database EMBL as of the end of May, 1999 using BLASTN algorithm Version 2.0.6 [Sep-16-1998], set to the following running parameters: Unix running command: blastall -p blastn -d embldb -e 10 -G 0 -E 0 -r 1 -v 30 -b 30 -i queryseq -o results. Multiple alignments of redundant sequences were used to build up reliable consensus sequences. Based on similarity to known sequences from other plant species, the isolated polynucleotides of the present invention identified as SEQ ID NOS. 1-53 and 78-164 were putatively identified as encoding polypeptides having similarity to the polypeptides shown above in Table 1.

The isolated cDNA sequences were compared to sequences in the EMBL DNA database using the computer algorithm BLASTN. The corresponding predicted

polypeptide sequences were determined and were compared to sequences in the SwissProt database using the computer algorithm BLASTP. Comparisons of DNA sequences provided in SEQ ID NOS: 78-164, to sequences in the EMBL DNA database (using BLASTN) and amino acid sequences provided in SEQ ID NOS: 165-304 to sequences in the SwissProt database (using BLASTP) were made as of May, 1999. Analysis of six-frame translations of the polynucleotides of SEQ ID NOS: 78-164, were also compared to and aligned with the six-frame translations of polynucleotides in the EMBL database using the TBLASTX program.

10 BLASTN Polynucleotide Analysis

The cDNA sequences of SEQ ID NOS: 1, 2, 4-6, 8-12, 15, 19, 21-23, 27-33, 35, 37-42, 44, 46-52, 78-80, 82, 83, 86, 89-92, 96-100, 104-113, 115, 117, 120, 122-130, 132-136, 138-158, 160, 163 and 164, were determined to have less than 40% identity, determined as described above, to sequences in the EMBL database using the computer algorithm BLASTN, as described above. The cDNA sequences of SEQ ID NOS: 3, 7, 14, 18, 20, 25, 34, 36, 53, 84, 85, 87, 88, 101, 114, 116, 118, 119, 131, 137, 159, 161 and 162 were determined to have less than 60% identity, determined as described above, to sequences in the EMBL database using BLASTN, as described above. The cDNA sequences of SEQ ID NOS: 16, 17, 26, 43, 45, 93, 94 and 121, were determined to have less than 75% identity, determined as described above, to sequences in the EMBL database using BLASTN, as described above. The cDNA sequences of SEQ ID NOS: 13, 24, 95, 102 and 103 were determined to have less than 90% identity, determined as described above, to sequences in the EMBL database using BLASTN, as described above.

25 BLASTP Amino Acid Analysis

The predicted amino acid sequences of SEQ ID NOS: 194-200, 202, 216, 223, 230, 235, 239, 240, 243, 250, 255, 259, 260, 263, 270, 272, 274, 278, 291, 292, 293, 296, 303 and 304 were determined to have less than 50% identity, determined as described above, to sequences in the SwissProt database using the BLASTP computer algorithm as described above. The predicted amino acid sequences of SEQ ID NOS: 166, 168-177, 179, 183-188, 192, 203-205, 207, 209-213, 218, 219, 221, 224, 225, 227-229, 231, 232, 234, 237, 242, 244, 245, 251, 253, 262, 267, 268, 269, 273, 276, 277, 279, 281, 282, 284, 286, 289, 290, 294, 295, 297, 298, 299, 300, 301 and 302 were determined to have less

than 75% identity, determined as described above, to sequences in the SwissProt database using the computer algorithm BLASTP, as described above. The predicted amino acid sequences of SEQ ID NOS: 165, 167, 178, 182, 189-191, 193, 201, 206, 208, 214, 215, 217, 220, 222, 226, 233, 238, 241, 246-250, 254, 256, 257, 258, 261, 264, 265, 266, 275, 5 280, 283, 285 and 288 were determined to have less than 90% identity, determined as described above, to sequences in the SwissProt database using the computer algorithm BLASTP, as described above. The predicted amino acid sequences of SEQ ID NOS: 180, 181 and 271, were determined to have less than 95% identity, determined as described above, to sequences in the SwissProt database using the computer algorithm BLASTP, as described above. 10 described above.

TBLASTX Analysis

The six-frame translations of the polynucleotide sequences of SEQ ID NOS: 78-164 were compared to and aligned with six-frame translations of polynucleotides in the EMBL database using the TBLASTX program version 2.0.6 [Sept-16-1998] set to the following running parameters: Unix running command: blastall -p blastn -d embldb -e 10 -G 0 -E 0 -v 30 -b 30 -i queryseq -o results. The translations of the polynucleotides of SEQ ID NOS: 82, 83, 90, 107-113, 115, 120, 122, 124-126, 129, 134-136, 142-144, 146-149, 152, 153, 155-158 and 164, were determined to have less than 50% identity, 15 determined as described above, to translations of polynucleotides in the EMBL database using the computer algorithm TBLASTX. The translations of the polynucleotides of SEQ ID NOS: 79, 81, 84-89, 91, 92, 96-101, 103, 105, 114, 116-118, 123, 131, 132, 137-141, 145, 150, 154 and 160-162, were determined to have less than 75% identity, determined as described above, to translations of polynucleotides in the EMBL database using the computer algorithm TBLASTX. 20 The translations of the polynucleotide sequences of SEQ ID NOS: 78, 80, 93, 95, 102, 104, 106, 119, 121, 127, 128, 130, 133, 151, 159 and 163, were determined to have less than 90% identity, determined as described above, to translations of polynucleotides in the EMBL database using the computer algorithm TBLASTX. 25 The translations of the polynucleotide sequence of SEQ ID NO: 94 was determined to have less than 95% identity, determined as described above, to translations of polynucleotides in the EMBL database using the computer algorithm TBLASTX. 30

Example 2

Use of an O-methyltransferase (OMT) Gene to Modify Lignin Biosynthesis

5 Transformation of tobacco plants with a *Pinus radiata* OMT gene

– Genetic constructs comprising sense and anti-sense nucleotides containing a polynucleotide comprising the coding region of the enzyme O-methyltransferase (OMT) (SEQ ID NO: 54) from *Pinus radiata* were constructed and inserted into *Agrobacterium tumefaciens* by direct transformation using published methods (An *et al.*, "Binary
10 vectors," in Gelvin SB and Schilperoort RA, eds., *Plant Molecular Biology Manual*, Kluwer Academic Publishers: Dordrecht, 1988). General methods for plant transformation are described in Horsch *et al.*, *Science* 227:1229-1231, 1985. The constructs of sense DNA were made by first cloning the PBK-CMV cDNA inserts into pART7 vectors. The pART7 vectors were then cut by restriction endonuclease *NotI* to remove the 35S-Insert-
15 OCS 3'UTR construct for cloning into the plant expression vector pART27 (Gleave A, *Plant Mol. Biol.* 20:1203-1207, 1992). The presence and integrity of the transgenic constructs were verified by restriction digestion and DNA sequencing.

Tobacco (*Nicotiana tabacum* cv. Samsun) leaf sections were transformed with the sense and anti-sense OMT constructs using the method of Horsch *et al.*, *Science*
20 227:1229-1231, 1985. Five independent transformed plant lines were established for the sense construct and eight independent transformed plant lines were established for the anti-sense construct for OMT. Transformed plants containing the appropriate gene construct were verified using Southern blot experiments. A "+" in the column labeled "Southern" in Table 2 below indicates that the transformed plant lines were confirmed as
25 independent transformed lines.

Expression of *Pinus* OMT in transformed plants

Total RNA was isolated from each independent transformed plant line created with the OMT sense and anti-sense constructs. The RNA samples were analyzed in Northern
30 blot experiments to determine the level of expression of the transgene in each transformed line.

The data shown in the column labeled "Northern" in Table 1 shows that the transformed plant lines containing the sense and anti-sense constructs for OMT all exhibited high levels of expression, relative to the background on the Northern blots.

OMT expression in sense plant line number 2 was not measured because the RNA sample showed signs of degradation. There was no detectable hybridization to RNA samples from empty vector-transformed control plants.

5 Modulation of OMT enzyme activity in transformed plants

The total activity of OMT enzyme, encoded by the *Pinus* OMT gene and by the endogenous tobacco OMT gene, was analyzed for each transformed plant line created with the OMT sense and anti-sense constructs. Crude protein extracts were prepared from each transformed plant and assayed using the method of Zhang *et al.*, *Plant Physiol.* 113:65-74,
10 1997. The data contained in the column labeled "Enzyme" in Table 2 shows that the transformed plant lines containing the OMT sense construct generally had elevated OMT enzyme activity, with a maximum of 199%, whereas the transformed plant lines containing the OMT anti-sense construct generally had reduced OMT enzyme activity, with a minimum of 35%, relative to empty vector-transformed control plants. OMT
15 enzyme activity was not estimated in sense plant line number 3.

Effects of *Pinus* OMT on lignin concentration in transformed plants

OMT is an enzyme involved in the biosynthesis of lignin. The concentration of lignin in the transformed tobacco plants was determined using the well-established
20 procedure of thioglycolic acid extraction (Freudenberg *et al.*, *Constitution and Biosynthesis of Lignin*, Springer-Verlag: Berlin, 1968). Briefly, whole tobacco plants, of an average age of 38 days, were frozen in liquid nitrogen and ground to a fine powder in a mortar and pestle. 100 mg of frozen powder from one empty vector-transformed control plant line, the five independent transformed plant lines containing the sense construct for
25 OMT and the eight independent transformed plant lines containing the anti-sense construct for OMT were extracted individually with methanol, followed by 10% thioglycolic acid and finally dissolved in 1 M NaOH. The final extracts were assayed for absorbance at 280 nm. The data shown in the column labeled "TGA" in Table 2 shows that the transformed plant lines containing the sense and the anti-sense OMT gene constructs all exhibited
30 significantly decreased levels of lignin, relative to the empty vector-transformed control plant lines.

TABLE 2

	plant line	transgene	orientation	Southern	Northern	Enzyme	TGA
5	1	control	na	+	blank	100	104
	1	OMT	sense	+	2.9E+6	86	55
	2	OMT	sense	+	na	162	58
	3	OMT	sense	+	4.1E+6	na	63
	4	OMT	sense	+	2.3E+6	142	66
10	5	OMT	sense	+	3.6E+5	199	75
	1	OMT	anti-sense	+	1.6E+4	189	66
	2	OMT	anti-sense	+	5.7E+3	35	70
	3	OMT	anti-sense	+	8.0E+3	105	73
	4	OMT	anti-sense	+	1.4E+4	109	74
15	5	OMT	anti-sense	+	2.5E+4	87	78
	6	OMT	anti-sense	+	2.5E+4	58	84
	7	OMT	anti-sense	+	2.5E+4	97	92
	8	OMT	anti-sense	+	1.1E+4	151	94

20 These data clearly demonstrate that polynucleotides identified from isolated cDNA obtained as in Example 1 and encoding polypeptides, may be assembled in DNA constructs and used to transform plants. The data furthermore demonstrates that transformed plants comprising genetic constructs exhibit varied levels of such enzyme expression and activity, and that the modulation of the metabolism of such an enzyme, manipulated by either sense or anti-sense expression of a gene encoding the enzyme, such as OMT, affects end product concentrations, such as the lignin concentration in the transformed plants.

Example 3

Use of a 4-Coumarate:CoA ligase (4CL) Gene to Modify Lignin Biosynthesis

Transformation of tobacco plants with a *Pinus radiata* 4CL gene

35 Sense and anti-sense constructs containing a DNA sequence including the coding region of 4CL (SEQ ID NO: 55) from *Pinus radiata* were inserted into *Agrobacterium tumefaciens* LBA4301 by direct transformation as described above in Example 2. The presence and integrity of the transgenic constructs were verified by restriction digestion and DNA sequencing.

40 Tobacco (*Nicotiana tabacum* cv. Samsun) leaf sections were transformed as described above. Five independent transformed plant lines were established for the sense

construct and eight independent transformed plant lines were established for the anti-sense construct for 4CL. Transformed plants containing the appropriate lignin gene construct were verified using Southern blot experiments. A "+" in the column labeled "Southern" in Table 3 indicates that the transformed plant lines listed were confirmed as independent
5 transformed lines.

Expression of *Pinus* 4CL in transformed plants

Total RNA was isolated from each independent transformed plant line created with the 4CL sense and anti-sense constructs. The RNA samples were analyzed in Northern blot experiments to determine the level of expression of the transgene in each transformed
10 line. The data shown in the column labeled "Northern" in Table 3 below shows that the transformed plant lines containing the sense and anti-sense constructs for 4CL all exhibit high levels of expression, relative to the background on the Northern blots. 4CL expression in anti-sense plant line number 1 was not measured because the RNA was not
15 available at the time of the experiment. There was no detectable hybridization to RNA samples from empty vector-transformed control plants.

Modulation of 4CL enzyme activity in transformed plants

The total activity of 4CL enzyme, encoded by the *Pinus* 4CL gene and by the endogenous tobacco 4CL gene in transformed tobacco plants, was analyzed for each
20 transformed plant line created with the 4CL sense and anti-sense constructs. Crude protein extracts were prepared from each transformed plant and assayed using the method of Zhang *et al.*, *Plant Physiol.* 113:65-74, 1997. The data contained in the column labeled "Enzyme" in Table 2 shows that the transformed plant lines containing the 4CL sense
25 construct had elevated 4CL enzyme activity, with a maximum of 258%, and the transformed plant lines containing the 4CL anti-sense construct had reduced 4CL enzyme activity, with a minimum of 59%, relative to empty vector-transformed control plants.

Effects of *Pinus* 4CL on lignin concentration in transformed plants

30 The concentration of lignin in samples of transformed plant material was determined as described in Example 2. The data shown in the column labeled "TGA" in Table 3, below, shows that the transformed plant lines containing the sense and the anti-sense 4CL gene constructs all exhibited significantly decreased levels of lignin, relative to

the empty vector-transformed control plant lines. These data demonstrate that the polynucleotides identified from isolated cDNA as obtained in Example 1 may be assembled into DNA constructs and used to transform plants. Transformed plants comprising such genetic constructs exhibit modified levels of enzyme expression and activity. The metabolism of the biosynthetic pathway involving the enzyme is also affected.

TABLE 3

	plant line	transgene	orientation	Southern	Northern	Enzyme	TGA
10	1	control	na	+	blank	100	92
	2	control	na	+	blank	100	104
	1	4CL	sense	+	2.3E+4	169	64
15	2	4CL	sense	+	4.5E+4	258	73
	3	4CL	sense	+	3.1E+4	174	77
	4	4CL	sense	+	1.7E+4	164	80
	5	4CL	sense	+	1.6E+4	184	92
	1	4CL	anti-sense	+	na	59	75
20	2	4CL	anti-sense	+	1.0E+4	70	75
	3	4CL	anti-sense	+	9.6E+3	81	80
	4	4CL	anti-sense	+	1.2E+4	90	83
	5	4CL	anti-sense	+	4.7E+3	101	88
	6	4CL	anti-sense	+	3.9E+3	116	89
25	7	4CL	anti-sense	+	1.8E+3	125	94
	8	4CL	anti-sense	+	1.7E+4	106	97

Example 430 Transformation of Tobacco using Lignin Biosynthetic Genes

Sense and anti-sense constructs containing DNA sequences including the coding regions of coumarate 3-hydroxylase (C3H) (SEQ ID NO: 56), ferulate-5-hydroxylase (F5H) (SEQ ID NO: 57), cinnamoyl-CoA reductase (CCR) (SEQ ID NO: 58) and coniferyl glycosyl transferase (CGT) (SEQ ID NO: 59) from *Eucalyptus grandis*, and phenylalanine ammonia-lyase (PAL) (SEQ ID NOS: 60 and 61), cinnamate 4-hydroxylase (C4H) (SEQ ID NOS: 62 and 63), phenolase (PNL) (SEQ ID NO: 64) and laccase (LAC) (SEQ ID NO: 65) from *Pinus radiata* were inserted into *Agrobacterium tumefaciens* LBA4301 by direct transformation as described above. The presence and

integrity of the transgenic constructs were verified by restriction digestion and DNA sequencing.

5 Tobacco (*Nicotiana tabacum* cv. Samsun) leaf sections were transformed as described in Example 2. Up to twelve independent transformed plant lines were established for each sense construct and each anti-sense construct listed in the preceding paragraph. Transformed plants containing the appropriate lignin gene construct were verified using Southern blot experiments. All of the transformed plant lines analyzed were confirmed as independent transformed lines. This demonstrates that transgenic plants with an expressed novel gene can be made, starting the whole process from an isolated cDNA
10 obtained as in Example 1.

Example 5

Manipulation of Lignin Content in Transformed Plants

15

Determination of transgene expression by Northern blot experiments

Total RNA was isolated from each independent transformed plant line described in Example 4. The RNA samples were analyzed in Northern blot experiments to determine
20 the level of expression of the transgene in each transformed line. The column labeled "Northern" in Table 4 shows the level of transgene expression for all plant lines assayed, relative to the background on the Northern blots. There was no detectable hybridization to RNA samples from empty vector-transformed control plants.

25 Determination of lignin concentration in transformed plants

The concentration of lignin in empty vector-transformed control plant lines and in up to twelve independent transformed lines for each sense construct and each anti-sense construct described in Example 5 was determined as described in Example 3. The column labeled "TGA" in Table 3 shows the thioglycolic acid extractable lignins for all plant lines
30 assayed, expressed as the average percentage of TGA extractable lignins in transformed plants versus control plants. The range of variation is shown in parentheses.

TABLE 4

	transgene	orientation	no. of lines	Northern	TGA
5	control	na	3	blank	100 (92-104)
	C3H	sense	5	3.7E+4	74 (67-85)
	F5H	sense	10	5.8E+4	70 (63-79)
	F5H	anti-sense	9	5.8E+4	73 (35-93)
10	CCR	sense	1	na	74
	CCR	anti-sense	2	na	74 (62-86)
	transgene	orientation	no. of lines	Northern	TGA
	PAL	sense	5	1.9E+5	77 (71-86)
	PAL	anti-sense	4	1.5E+4	62 (37-77)
15	C4H	anti-sense	10	5.8E+4	86 (52-113)
	PNL	anti-sense	6	1.2E+4	88 (70-114)
	LAC	sense	5	1.7E+5	na
	LAC	anti-sense	12	1.7E+5	88 (73-114)

20 Transformed plant lines containing the sense and the anti-sense lignin biosynthetic gene constructs all exhibited significantly decreased levels of lignin, relative to the empty vector-transformed control plant lines. The most dramatic effects on lignin concentration were seen in the F5H anti-sense plants with as little as 35% of the amount of lignin in control plants, and in the PAL anti-sense plants with as little as 37% of the amount of

25 lignin in control plants. These data clearly indicate that the concentration of a polynucleotide, such as lignin, as measured by the TGA assay, can be directly manipulated by conventional anti-sense methodology and also by sense over-expression using the inventive lignin biosynthetic genes, starting the whole process from an isolated cDNA obtained as in Example 1.

30

Example 6

Modulation of Lignin Enzyme Activity in Transformed Plants

35

The activities and substrate specificities of selected lignin biosynthetic enzymes were assayed in crude extracts from transformed tobacco plants containing sense and anti-sense constructs for PAL (SEQ ID NO: 60), PNL (SEQ ID NO: 64) and LAC (SEQ ID NO: 65) from *Pinus radiata*, and CGT (SEQ ID NO: 59) from *Eucalyptus grandis*.

40

Enzyme assays were performed using published methods for PAL (Southerton SG and Deverall BJ, *Plant Path.* 39:223-230, 1990); CGT (Vellekoop *et al.*, *FEBS Lett.*

330:36-40, 1993); PNL (Espin *et al.*, *Phytochemistry* 44:17-22, 1997); and LAC (Bao *et al.*, *Science* 260:672-674, 1993). The data shown in the column labelled "Enzyme" in Table 5 shows the average enzyme activity from replicate measures for all plant lines assayed, expressed as a percent of enzyme activity in empty vector-transformed control plants. The range of variation is shown in parentheses.

TABLE 5

10	<u>Transgene orientation</u>	<u>no. of lines</u>	<u>enzyme</u>
	control na	3	100
	PAL sense	5	87 (60-124)
	PAL anti-sense	3	53 (38-80)
	CGT anti-sense	1	89
15	PNL anti-sense	6	144 (41-279)
	LAC sense	5	78 (16-240)
	LAC anti-sense	11	64 (14-106)

20 All of the transformed plant lines, except the PNL anti-sense transformed plant lines, showed average enzyme activities that were significantly lower than the activities observed in empty vector-transformed control plants. The most dramatic effects on lignin enzyme activities were seen in the PAL anti-sense transformed plant lines, in which all of the lines showed reduced PAL activity, and in the LAC anti-sense transformed plant lines, which showed as little as 14% of the LAC activity in empty vector-transformed control plant lines. These results demonstrate that enzyme activity can be modulated by transforming plants with polynucleotides encoding an enzyme of interest, starting the whole process from polynucleotides encoding enzymes of interest isolated from cDNA as described in Example 1.

Example 7

Functional Identification of Lignin Biosynthetic Genes

35 Sense constructs containing DNA sequences including the coding regions for PAL (SEQ ID NO: 61), OMT (SEQ ID NO: 54), 4CL (SEQ ID NOS: 55 and 66) and POX (SEQ ID NO: 67) from *Pinus radiata*, and OMT (SEQ ID NOS: 68 and 69), CCR (SEQ ID NOS: 70 - 72), CGT (SEQ ID NOS: 59 and 73) and POX (SEQ ID NOS: 74 and 75)

from *Eucalyptus grandis* were inserted into the commercially available protein expression vector, pProEX-HT (Gibco BRL). The resultant constructs were transformed into *E. coli* XL1-Blue (Stratagene), which were then induced to produce recombinant protein by the addition of IPTG. Purified proteins were produced for the *Pinus* OMT and 4CL constructs
 5 and the *Eucalyptus* OMT and POX constructs using Ni column chromatography (Janknecht *et al.*, *Proc. Natl. Acad. Sci. USA* 88:8972-8976, 1991). Enzyme assays for each of the purified proteins conclusively demonstrated the expected substrate specificity and enzymatic activity for the genes tested.

The data for two representative enzyme assay experiments, demonstrating the
 10 verification of the enzymatic activity of a *Pinus radiata* 4CL gene (SEQ ID NO: 55) and a *Pinus radiata* OMT gene (SEQ ID NO: 54), are shown below in Table 6. For the 4CL enzyme, one unit equals the quantity of protein required to convert the substrate into product at the rate of 0.1 absorbance units per minute. For the OMT enzyme, one unit equals the quantity of protein required to convert 1 pmole of substrate to product per
 15 minute.

TABLE 6

20	transgene	purification step	total ml extract	total mg protein	total units activity	% yield activity	fold purification
	4CL	crude	10 ml	51 mg	4200	100	1
		Ni column	4 ml	0.84 mg	3680	88	53
25	OMT	crude	10 ml	74 mg	4600	100	1
		Ni column	4 ml	1.2 mg	4487	98	60

The data shown in Table 6 demonstrate that both the purified 4CL enzyme and the
 30 purified OMT enzyme show high activity in enzyme assays, confirming the identification of the 4CL and OMT genes. Crude protein preparations from *E. coli* transformed with empty vector show no activity in either the 4CL or the OMT enzyme assay. This demonstrates that the function of an isolated novel cDNA with only a putative function can be confirmed, starting the whole process from an isolated cDNA obtained as in
 35 Example 1.

Example 8

Demonstration of the Presence / Absence of Unique Sequence Identifiers in Plants

5 Transgenic tobacco plants were created using unique identifier sequences which are not found in tobacco. The unique identifier sequences inserted were isolated from *Pinus radiata*, SEQ ID NO: 76, and *Eucalyptus grandis*, SEQ ID NO: 77. The unique identifier sequences were inserted into *Agrobacterium tumefaciens* LBA4301 (provided as a gift by Dr. C. Kado, University of California, Davis, CA) by direct transformation using
10 published methods (An *et al.*, "Binary vectors," in Gelvin SB and Schilperoort RA, eds., *Plant Molecular Biology Manual*, Kluwer Academic Publishers: Dordrecht, 1988). The presence and integrity of the unique identifier sequences in the *Agrobacterium* transgenic constructs were verified by restriction digestion and DNA sequencing.

Tobacco (*Nicotiana tabacum* cv. Samsun) leaf sections were transformed using the
15 method of Horsch *et al.*, *Science* 227:1229-1231, 1985. Three independent transformed plant lines were established for each unique sequence identifier used. Two empty-vector control plant lines were established using an empty gene transfer vector that lacked a unique sequence identifier.

The uniqueness of the sequence identifiers was assayed using Southern blot
20 analyses to test for the presence of the sequence identifier in the genome of the plants. If the sequence identifier is unique and therefore useful as a tag, then the sequence identifier should be clearly absent in plants which have not been tagged and it should be clearly present in plants which have been tagged. In the present example, the unique identifiers would be expected to be absent in the empty-vector transformed control plants. The
25 unique identifier would be expected to be present in the transgenic plants transformed with the unique sequence identifiers.

Genomic DNA was prepared from empty-vector transformed control plants and plants transformed with unique sequence identifiers using the cetyltrimethyl-ammonium bromide (CTAB) extraction method of Murray MG and Thompson WF, *Nucleic Acids*
30 *Res.* 8:4321-4325, 1980. The DNA samples were digested with the restriction enzyme *EcoRI* in the case of the plants transformed with the *Pinus* unique sequence identifier (SEQ ID NO: 76) and the restriction enzyme *XbaI* in the case of the plants transformed with the *Eucalyptus* unique sequence identifier (SEQ ID NO: 77). The DNA fragments

produced in the restriction digests were resolved on a 1% agarose gel; the left panel of Fig. 2 and the right panel of Fig. 2 show the DNA fragment patterns of the DNA samples from the *Pinus* and *Eucalyptus* experiments, respectively.

After the agarose gel electrophoresis step, the DNA samples were transferred to
5 Hybond-N+ nylon membranes (Amersham Life Science, Little Chalfont, Buckinghamshire, England) using methods established by Southern, *J. Mol. Biol.* 98:503-517, 1975. The nylon membranes were probed with radioactively-labeled probes for the unique sequence identifiers identified above and washed at high stringency (final wash: 0.5 X salt sodium citrate buffer (SSC) plus 0.1% sodium dodecyl sulfate (SDS), 15
10 minutes at 65°C). The hybridization of the probes to complementary sequences in the genomic DNA samples was detected using auto-radiography.

The results are shown in Figs. 3 and 4.

Fig. 3 (corresponding to the left panel of Fig. 2) shows the hybridization pattern detected in the Southern blot analysis using a probe derived from the *Pinus* sequence
15 identifier (SEQ ID NO: 76). Lanes A-B contain DNA samples from empty-vector transformed control plants and Lanes C-E contain DNA from plants transformed with SEQ ID NO: 76. There is no hybridization in Lanes A-B indicating that SEQ ID NO: 76 is not present in empty-vector transformed tobacco plants; that is, SEQ ID NO: 76 is a unique tag suitable for unambiguous marking of tobacco plants. There is strong
20 hybridization in Lanes C-E, indicating that the plants which received SEQ ID NO: 76 via transformation have been clearly and unambiguously tagged with the unique sequence contained in SEQ ID NO: 76.

Fig. 4 (corresponding to the right panel of Fig. 2) shows the hybridization pattern detected in the Southern blot analysis using a probe derived from the *Eucalyptus* sequence
25 identifier (SEQ ID NO: 77). Lanes A-B contain DNA samples from empty-vector transformed control plants and Lanes C-E contain DNA from plants transformed with SEQ ID NO: 77. There is no hybridization in Lanes A-B indicating that SEQ ID NO: 77 is not present in empty-vector transformed tobacco plants; that is, SEQ ID NO: 77 is a unique tag suitable for unambiguous marking of tobacco plants. There is strong
30 hybridization in Lanes C-E indicating that the plants which received SEQ ID NO: 77 via transformation have been clearly and unambiguously tagged with the unique sequence contained in SEQ ID NO: 77.

The data clearly demonstrates the utility of the sequences disclosed in this specification for the purposes of unambiguously tagging transgenic materials. A unique sequence was selected from a large number of potential tags and shown to be absent in the genome of the organism to be tagged. The tag was inserted into the genome of the
5 _ organism to be tagged and a well-established DNA detection method was used to clearly detect the unique sequence identifier used as the tag.

Because of the sequence-specific detection methods used in the example, a user of the invention disclosed in this specification has both a high likelihood of finding a sequence identifier, among the list which has been disclosed, which will be useful for
10 tagging any given organism and an unequivocal method for demonstrating that a tagged organism could only have acquired a given tag through the deliberate addition of the unique sequence to the genome of the organism to be tagged. If the user of this invention maintains the precise sequence of the tag used in a given organism as a secret, then any disputes as to the origin and history of the organism can be unambiguously resolved using
15 the tag detection techniques demonstrated in the present example.

SEQ ID NOS: 1-304 are set out in the attached Sequence Listing. The codes for nucleotide sequences used in the attached Sequence Listing, including the symbol "n," conform to WIPO Standard ST.25 (1998), Appendix 2, Table 1.

All references cited herein, including patent references and non-patent
20 publications, are hereby incorporated by reference in their entirety. While in the foregoing specification, this invention has been described in relation to certain preferred embodiments, and many details have been set forth for purposes of illustration, it will be apparent to those skilled in the art that the invention is susceptible to additional embodiments and that certain of the details described herein may be varied considerably
25 without departing from the basic principles of the invention.

Claims:

1. An isolated polynucleotide comprising a polynucleotide sequence selected from the group consisting of: (1) the sequences recited in SEQ ID NOS: 1-53 and 78-164; (2) complements of the sequences recited in SEQ ID NOS: 1-53 and 78-164; (3) reverse complements of the sequences recited in SEQ ID NOS: 1-53 and 78-164; (4) reverse sequences of the sequences recited in SEQ ID NOS: 1-53 and 78-164; (5) sequences comprising a polynucleotide sequence having at least 40% identity to a compare sequence selected from the polynucleotide sequences recited in SEQ ID NOS: 1, 2, 4-6, 8-12, 19, 21-23, 28-33, 35, 37-42, 44, 46-52, 78-80, 82, 83, 86, 89-92, 96-100, 104-113, 115, 117, 120, 122-130, 132-136, 138-158, 160, 163 and 164, the percentage identity determined by aligning the sequence and the compare sequences using the BLASTN algorithm version 2.04 set at the parameter values described herein, identifying the number of identical nucleic acids over aligned portions of the sequence and the compare sequences, dividing the number of identical nucleic acids by the total number of nucleic acids of the compare sequence, and multiplying by 100 to determine the percentage identity; (6) sequences comprising a polynucleotide sequence having at least 60% identity to a compare sequence selected from the polynucleotide sequences recited in SEQ ID NOS: 3, 7, 14, 18, 20, 25, 34, 36, 53, 84, 85, 87, 88, 101, 114, 116, 118, 119, 131, 137, 159, 161 and 162, the percentage identity determined as described in (5) above; (7) sequences comprising a polynucleotide sequence having at least 75% identity to a compare sequence selected from the polynucleotide sequences recited in SEQ ID NOS: 16, 17, 26, 43, 45, 93, 94 and 121, the percentage identity determined as described in (5) above; (8) sequences comprising a polynucleotide sequence having at least 90% identity to a compare sequence selected from the nucleotide sequences recited in SEQ ID NOS: 13, 24, 95, 102 and 103, the percentage identity determined as described in (5) above; (9) sequences comprising a polynucleotide sequence that hybridizes to a polynucleotide comprising a sequence recited in (1) – (8) above under stringent hybridization conditions; (10) sequences comprising a polynucleotide sequence that is a 100-mer of a sequence recited in (1) – (8) above; (11) sequences comprising a polynucleotide sequence that is a 40-mer of a sequence recited in (1) – (8) above; and (12) sequences

5 2. An isolated oligonucleotide probe or primer comprising at least 10 contiguous residues complementary to 10 contiguous residues of a nucleotide sequence recited in Claim 1.

4. A transgenic cell comprising a genetic construct according to claim 3.

6. A genetic construct comprising, in the 5'-3' direction:

(c) a gene termination sequence.

7. The construct of claim 6 wherein the polynucleotide is in a sense orientation.

8. The construct of claim 6 wherein the polynucleotide is in an antisense orientation.

25 9. The construct of claim 6 wherein the gene promoter sequence and gene termination sequences are functional in a plant host.

10. A transgenic cell comprising a construct of claim 6.

11. The transgenic cell of claim 10 wherein the polynucleotide is in a sense orientation.

30 12. The transgenic plant cell of claim 10 wherein the polynucleotide is in an antisense orientation.

13. A transgenic cell according to claim 10, wherein the cell is selected from one of the following: a bacterial cell; an insect cell; a yeast cell; a mammalian cell; and a plant cell.
14. A plant comprising a transgenic cell according to claim 9, or fruit or seeds or progeny thereof.
15. The plant of claim 14 wherein the plant is a woody plant.
16. The plant of claim 15 wherein the plant is selected from the group consisting of eucalyptus and pine species.
17. A method for modulating one or more of the content, the composition, and the metabolism of an enzyme involved in an isoprenoid biosynthetic pathway in an organism, comprising stably incorporating into the genome of the organism a construct of claim 3.
18. A method according to claim 17, wherein the organism is a plant.
19. A method for modulating one or more of the content, the composition, and the metabolism of an isoprenoid compound in an organism comprising stably incorporating into the genome of the organism a construct of claim 6.
20. A method according to claim 19, wherein the organism is a plant.
21. A method for producing an organism having one or more of altered isoprenoid content, altered isoprenoid composition and altered isoprenoid metabolism, comprising:
- (a) transforming a host cell with a construct of claim 3 to provide a transgenic host cell; and
 - (b) cultivating the transgenic host cell under conditions conducive to growth and regeneration.
22. A method according to claim 21, wherein the organism is a plant and the host cell is a plant cell.
23. An isolated polypeptide encoded by a polynucleotide of claim 1.
24. A polypeptide of claim 23 having enzymatic activity in an isoprenoid biosynthetic pathway in a plant.
25. An isolated polypeptide comprising an amino acid sequence expressed from a polynucleotide that hybridizes to a nucleotide sequence set forth as SEQ ID NOS: 1-53 and 78-164 under stringent hybridization conditions.

26. An isolated polypeptide comprising a polypeptide sequence selected from the group consisting of: (1) the sequences set forth in SEQ ID NOS: 165-286 and 288-304; (2) sequences comprising a polypeptide sequence having at least 50% identity to a compare sequence selected from the polypeptide sequences recited in
5 SEQ ID NOS: 194-200, 202, 216, 223, 230, 235, 239, 240, 243, 250, 255, 259, 260, 263, 270, 272, 274, 278, 291, 292, 293, 296, 303 and 304; (3) sequences comprising a polypeptide sequence having at least 75% identity to a compare sequence selected from the polypeptide sequences recited in SEQ ID NOS: 166, 168-177, 179, 183-188, 192, 203-205, 207, 209-213, 218, 219, 221, 224, 225, 227-
10 229, 231, 232, 234, 237, 242, 244, 245, 251, 253, 262, 267, 268, 269, 273, 276, 277, 279, 281, 282, 284, 286, 289, 290, 294, 295, and 297-302; (4) sequences comprising a polypeptide sequence having at least 90% identity to a compare sequence selected from the polypeptide sequences recited in SEQ ID NOS: 165, 167, 178, 182, 189-191, 193, 201, 206, 208, 214, 215, 217, 220, 222, 226, 233,
15 238, 241, 246-250, 254, 256-258, 261, 264, 265, 266, 275, 280, 283, 285 and 288; (5) sequences comprising a polypeptide sequence having at least 95% identity to a compare sequence selected from the polypeptide sequences recited in SEQ ID NOS: 180, 181 and 271; (6) sequences comprising a polypeptide sequence that is a 100-mer of a sequence recited in (1) – (5) above having at least 100 residues; (7)
20 sequences comprising a polypeptide sequence that is a 40-mer of a sequence recited in (1) – (5) above having at least 40 residues; and (8) sequences comprising a polypeptide sequence that is a 20-mer of a sequence recited in (1) – (5) above.
27. A method for modulating one or more of the content, the composition and the metabolism of an isoprenoid compound in an organism, comprising administering
25 an isolated polypeptide of claim 26 to the organism.
28. A method according to claim 27, wherein the organism is a plant, and administration of the isolated polypeptide is topical.
29. A method according to claim 27, wherein the organism is a mammal, and administration of the isolated polypeptide is systemic.

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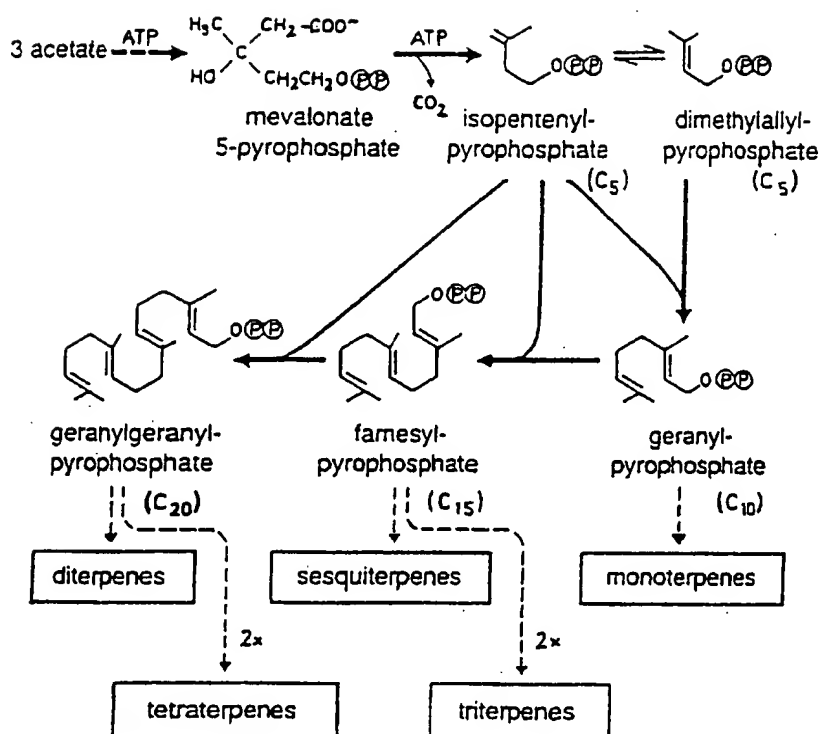


FIGURE 1

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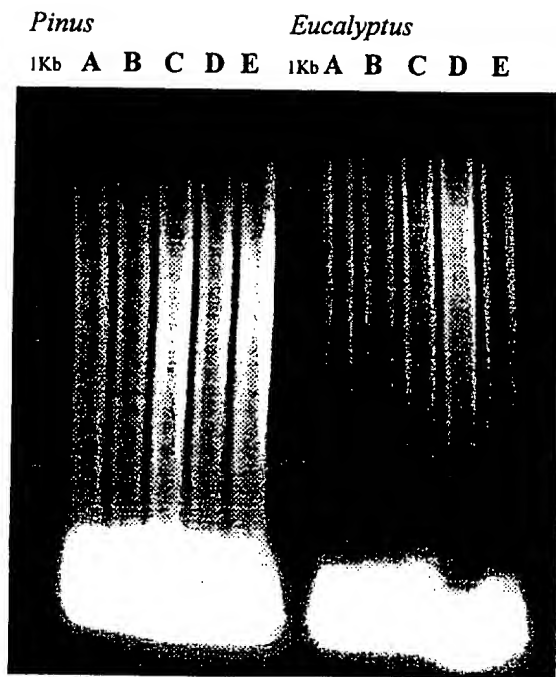


FIGURE 2

3/4

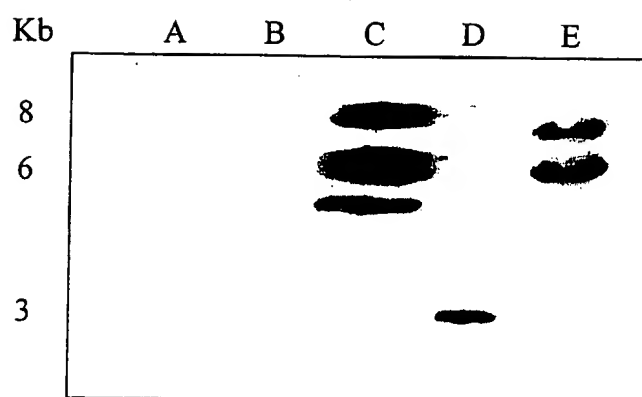


FIGURE 3

4/4

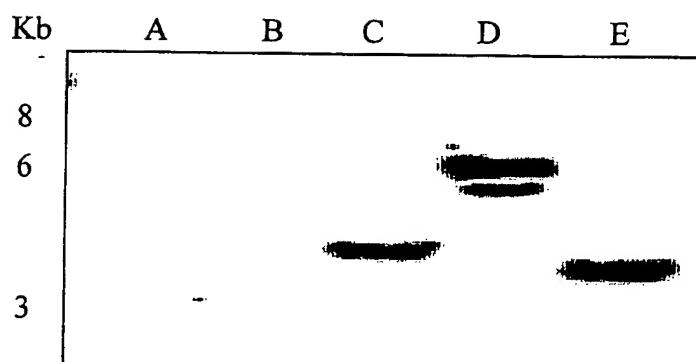


FIGURE 4

SEQUENCE LISTING

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of Isoprenoid Content, Compositition and Metabolism

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<210> 4
 <211> 371
 <212> DNA
 <213> Eucalyptus grandis

<400> 4						
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tgattcagca	aatgtcgcaa	aagtgaaggt	cctctaccac	gagatcaatc	ttcagggata	300
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<210> 5
 <211> 408
 <212> DNA
 <213> Pinus radiata

<400> 5						
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aaggaagatt	tatcttgctt	cgacccaaag	aaggccgcac	cgttgttggg	tattgcagag	360
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<210> 6
 <211> 590
 <212> DNA
 <213> Eucalyptus grandis

<400> 6						
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<210> 7

<211> 699

<212> DNA

<213> Eucalyptus grandis

<400> 7

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ctgctttcct	ggagctgggg	aaaagttacc	aagaggcaat	tgatgatatt	actaaaagaa	660
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<210> 8

<211> 373

<212> DNA

<213> Pinus radiata

<400> 8

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gtggcgattg	cggtagccat	tggctttgtt	tctgtattat	tgtcgtatta	tatagttttg	180
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tcaacagatg	acaatggcat	tgccatcgaa	gctgctggag	gaacggatgt	tatcatcgtg	300
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<210> 9

<211> 373

<212> DNA

<213> Pinus radiata

<400> 9

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gctttgtttc	tgtattattg	tcgtattata	tagttttgag	caggtggaag	cgcagatcca	240
acggattacg	gggaatacag	agcaaaaagt	tcgaaaagtc	aacagatgac	aatggcattg	300
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<210> 10

<211> 825

<212> DNA

<213> Eucalyptus grandis

<400> 10

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aaatagaagc	atgcctgccca	atctccaacc	cacccttgga	gctcttctga	tgggagacgc	120

gttcaacatg	cgccatccat	tgacaggagg	aggaatgacc	gtggetcttt	ctgatatcgt	180
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tgcagcaaa	ctcgataaag	gcatgaatat	gcgggtcaagt	tcgtttcttc	tagtcacatt	720
cctctctcgt	agtagatgga	gaggcctgcc	agttatactg	cactcggaag	agaattgtgc	780
attggaacta	tagatttcgt	tacgataagt	agattcattc	aagaa		825

<210> 11

<211> 394

<212> DNA

<213> Eucalyptus grandis

<400> 11

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tcgtcatctc	cggcctcgac	atcctcgacg	ccctcgatcg	cgtacacaaa	gatgcgggtg	360
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<210> 12

<211> 245

<212> DNA

<213> Eucalyptus grandis

<400> 12

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ggcacattgg	aagaacttgc	actcctcaat	gaagcaatcc	aaaagtggga	ttttgatgcc	120
atggatggat	taccagagta	tatgcaagct	tattttcaagg	agttttctcca	gctctatgaa	180
tatatgggga	atcaattggc	cgcaaaaagga	agatcgtacc	gccttatcta	cgcaaaaagaa	240
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<210> 13

<211> 375

<212> DNA

<213> Pinus radiata

<400> 13

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gagatagggg	acgtcctcga	gtatgggttg	cacacgtatt	tgccacgatt	ggaagcaagg	180
aattacatcg	acgtcctcgg	acaggacact	gaaaacagca	agtcatatat	gaagaccgag	240
aaactttctg	aacttgcaaa	gttggagttc	aacatctttc	acgccttaca	aaagcgagag	300
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<210> 14

<211> 824

<212> DNA

<213> Pinus radiata

<400> 14

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aagtccagct	agaagcatat	accaaagaag	cagaatggct	tgcagtgaga	tacgtgccat	180
cctatgatga	atatataggg	aacgcgagtg	tttcaatagc	attgggaaca	gtggttctta	240
tcagcgctct	ttttactggg	gagattctta	cagatgacat	actctccaaa	attggtcgcg	300
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<210> 15

<211> 1271

<212> DNA

<213> *Eucalyptus grandis*

<400> 15

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atcaagtcgg	tcgtcgaggg	aaagatgccg	tcgtactcgc	tcgagtccaa	gcttggggac	180
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<210> 16

<211> 372

<212> DNA

<213> *Pinus radiata*

<400> 16

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tcgtacggtg	agcatgctga	cagacttatt	ggggaagtga	aggggatttt	caactcattt	360
tcgattgcag	at					372

<210> 17

<211> 520

<212> DNA

<213> *Pinus radiata*

<400> 17

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aacgtttggg	gatcgacaga	catttccaaa	ctgaaataaa	agttgctctt	gactatgggt	480
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<210> 18

<211> 435

<212> DNA

<213> Pinus radiata

<400> 18

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ttacagttat	tggagcgaaa	aaggcattgg	atatgggaga	gagagcgcta	ttactgatct	420
caacacaact	tcctt					435

<210> 19

<211> 320

<212> DNA

<213> Pinus radiata

<400> 19

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gtcacatccc	ggatcactac					320

<210> 20

<211> 626

<212> DNA

<213> Eucalyptus grandis

<400> 20

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<210> 21

<211> 490

<212> DNA

<213> Eucalyptus grandis

<400> 21

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<210> 22

<211> 396

<212> DNA

<213> Eucalyptus grandis

<400> 22

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atccacaaaa	gttactctcg	acttgatcat	gaagatttta	aggtagatga	ccttcacaca	180
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gacaaattca	agatagcaac	gggaacttcc	gagagtcgac	tcataagtga	tgtgcgggga	300
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<210> 23

<211> 396

<212> DNA

<213> Eucalyptus grandis

<400> 23

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caaatccaac	gccttgggaat	tgagtaccat	tttgaacgtg	aaatagatga	gcaattagaa	360
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<210> 24

<211> 700

<212> DNA

<213> Pinus radiata

<400> 24

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gactcagctc	cattttctgc	tgctggaccg	ccttcttgcc	actctttctt	gtgttcttgc	480
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caatctatta	aaggatgaaa	gtgatgaaga	cagcttggtg	acagactttg	aggtcaactt	600
tccttttctg	ttgagagaag	ctcaatcttt	ccaacttgaa	ctcccttatg	acctgcctta	660
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<210> 25

<211> 1513

<212> DNA

<213> Pinus radiata

<400> 25

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<210> 26

<211> 295

<212> DNA

<213> Pinus radiata

<400> 26

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gagatgggtg	aaagaatcgg	gttcccctga	gctaaccctc	gctcgacatc	gttacgtgga	180
attctacact	ttgggtctgtg	gcattgacat	ggagcctaaa	gatgccgcat	tcagactgag	240
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<210> 27

<211> 191

<212> DNA

<213> Pinus radiata

<400> 27

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gaaatggcga	aagaagcaca	gaaatcacaa	ggccgagaca	cgctcggcta	tgttcgacag	120
gctgtaataa	caattgatat	gctatgcata	tattttgaata	aacaaatact	cgttggccat	180
ctattttatt	t					191

<210> 28

<211> 373

<212> DNA

<213> Pinus radiata

<400> 28

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gggtgcctgg	actttatcta	caagagatca	tgggtgggtt	gtggctgact	gctctgctga	180
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accccagcaa	aggcttttcg	cttgtgtcaa	ttatctactt	tccatgcaga	atacagacgg	300
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<210> 29

<211> 1411

<212> DNA

<213> *Eucalyptus grandis*

<400> 29

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accagatgga	cacggacaac	aaactcttca	atgtgggctg	cttgctcgtg	gccactctcg	180
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<210> 30

<211> 689

<212> DNA

<213> *Eucalyptus grandis*

<400> 30

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ccaaaaaagt	tgtaaagggt	acgcaaaaag	agcgaggatg	tggaagctca	agattggaga	180
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cctgagggag	aagaacttca	agcagacgat	ccctccgggt	aagatcgagg	atggcgagga	420
tatcacctat	gagaaggcga	cgcggccgtg	gaagcgggtg	gtcagcttct	ggtctactct	480
gcagtccagc	catggccact	ggcctgccga	gaacgcgggc	cccattgcgt	tctacttccc	540
tcccctgggt	atgagtctct	acgtgactgg	acatctgaac	aacgttttcc	atgccgagca	600
tcgcagggag	atcctccgct	acatttacta	ccatcagaac	gaagacgggtg	gctggggact	660
gcacattgaa	ggccacagca	cgatgatcg				689

<210> 31

<211> 393

<212> DNA

<213> *Pinus radiata*

<400> 31

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ccaaggcaat	ggagaacgaa	gatggagtgt	caggagcagt	aaaggcattt	cataaacatt	180
tacccaaaaa	aatgccacaa	ccactgccac	cacctacaga	tcatagtcta	attgattcct	240
tctttacagg	tggtggaaag	gtttttgggt	gtggctgatt	gtaccttaat	ataacgttta	300
tcaactatta	ggctgggatg	ctttgaagg	aaaaactgaa	aatcgagtgc	ttgaacttgt	360
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<210> 32

<211> 519

<212> DNA

<213> Pinus radiata

<400> 32

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gactggatct	gcactggctg	gtgccatggg	aggattcaat	gcccattgca	ccaatatcgt	180
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tattacgatg	atggaggcat	ccaacgatgg	aaaggatctt	catgtatcag	tcaccatgcc	300
ttgtattgag	gttggaacag	taggaggtgg	aacgcagttg	gcctcacaag	ctgcttgctt	360
gaatatgctc	ggagtgaag	gagcgaataa	ggaatcccc	ggagcgaatg	ctcagacttt	420
ggccagaatt	gtggcaggag	cagttttggc	tggagagctg	tctctcatgt	ctgccttagc	480
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<210> 33

<211> 302

<212> DNA

<213> Pinus radiata

<400> 33

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agaaagattc	actgaatcgc	ttactaaaac	tctgtttcag	gaatggcaac	aggaggagga	180
gcgttgatc	tgccctcagg	aatgggaggg	aacattgaga	aagaacaaat	gctgaccgct	240
gttgaagagt	acgaaaaata	tcacatgtac	tatggtgggtg	atgaaggctc	gagaaaatct	300
aa						302

<210> 34

<211> 508

<212> DNA

<213> Eucalyptus grandis

<400> 34

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tgccgacggg	ccttgccggg	aagatggaca	agagcgacgt	cctgtccgcc	gttgacaagt	180
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acatgggtgaa	taaatattat	gatcttgcta	ccagctttta	tgagttcggc	tggggagaaat	300
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gcgggaattg	tggaccgctt	agggaaatag	ctcgattcag	ctccgcactc	gttacaggat	480
taaacacaaa	tgagtaccag	ataacaag				508

<210> 35

<211> 353

<212> DNA

<213> Pinus radiata

<400> 35

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cagaataaagc	ttggtctatg	atccccacac	agatgaatac	tacaatgccc	ctgggggtgga	120
gactagagtg	ccttattttg	gttcaacaga	aggaatgaag	taccttgatc	cctgcttcaa	180
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cggtaaatcc	ctttttgggtg	caccctatga	tttccgttac	ggtcctggaa	caaagtccctc	300
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<210> 36

<211> 82

<212> DNA

<213> Eucalyptus grandis

<400> 36

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tgcttggtat	gggtcccatc	ca				82

<210> 37

<211> 474

<212> DNA

<213> Pinus radiata

<400> 37

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atggttgaag	actatggcaa	ctattggata	cacaggtggc	tacattgcaa	atggggctat	180
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ccaggacaca	tgattacatt	ctgggtgctg	gttgtgctgc	gccaagtggg	agcgattgaa	360
actcacagcg	gatatgactt	tccgtggact	cttaccaaat	taattccttt	ctatggaggc	420
gcggagtatc	atgactacca	tcattatgta	ggaggacaaa	gtcaaagcaa	cttt	474

<210> 38

<211> 340

<212> DNA

<213> Eucalyptus grandis

<400> 38

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caactactgg	ttgcacagat	tattgcactg	caaatggggg	tatgaaaaga	tccacagcgt	120
tcaccacgag	tacagcgctc	cgatcggttt	tgcggcgcca	tacgcgcact	gggctgaggt	180
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gttgtggttg	tggattgctt	tgcggcagat	tgaggccatc	gactactcac	agcgggtacg	300
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<210> 39

<211> 487

<212> DNA

<213> Eucalyptus grandis

<400> 39

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cgcgaagctc	ttctgttgat	gtgctagggt	ccctgcaatt	cttccgcccc	gattcctatc	420
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caaagat						487

<210> 40

<211> 571
 <212> DNA
 <213> Pinus radiata

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<210> 41
 <211> 512
 <212> DNA
 <213> Eucalyptus grandis

<400> 41
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<210> 42
 <211> 445
 <212> DNA
 <213> Pinus radiata

<400> 42
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 cgggtgtata ccccggaagc ccaaactctgc tcatcccgca ctactggga gcagaacttg 180
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 aaaatatgaa ttgcttcttc agcaa 445

<210> 43
 <211> 412
 <212> DNA
 <213> Eucalyptus grandis

<400> 43
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 cggcgctca tgttcgagga cgaatgcac ttggtggatg agaacgacaa tgcctgtcgt 180
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 gccacaaagg taacattccc ccttggtgtg acaaacacct gctgcagcca tccattgtac 360
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<210> 44
 <211> 834
 <212> DNA
 <213> Pinus radiata

<400> 44

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gattggaacc	gacatcgaag	acttcaagt	ctcttggtg	gtggtgcaag	cgcttgaacg	180
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aaaagggtca	tcacgattct	ttcagtgggt	agtccaacaa	ctctgatgga	gttcatectc	720
ggttacagca	tagtcatatc	tgaagtcgta	ctgtaggaaa	gtgttcacaa	cagatggcac	780
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<210> 45
 <211> 389
 <212> DNA
 <213> Pinus radiata

<400> 45

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gaacctaaat	tgatctcggc	agttcgggaat	tttcaaattct	caaattctcg	gaggggttcc	120
tgcgctttga	tcgttcgaga	tgggggaatc	tgaggagagt	ttgggtgcag	gttcgaatct	180
caagtcagct	gctgtgttgg	agcaggcaaa	gaaacacctt	gccacagacg	ctgcccaaga	240
cctcaagaag	aagatcggcc	ttgtctatca	gctcaacatt	tcaccaaga	aaattggaat	300
agctgaggag	gtgttcgttg	tggaacctcaa	gaatggcaaa	gtcactaaag	gaccatatga	360
aggaaagcca	gatgcaacat	tttcctttg				389

<210> 46
 <211> 469
 <212> DNA
 <213> Pinus radiata

<400> 46

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tttctccag	gtttcatgtt	tgggatagca	acttctgctt	atcagtgtga	aggagctgcc	120
aacgaagggtg	gaaaaggccc	aagcatctgg	gactcatttt	cacgaacacc	aggcaaaatt	180
cttgatggaa	gcaacgggtga	tgtagcagt	gatcagtatc	atcggtataa	ggaagatgta	240
aaactgatga	aagatatggg	agtggatacc	tacagattct	cgctttcatg	gcctcgtata	300
tttccaaagg	gaaaaggaga	gatcaatgag	gaaggagtag	cctattacaa	taacctcatc	360
aatgaactcc	tccagaatgg	aatccaagcg	tctgtcactt	tgtttctactg	ggatactccc	420
cagtctctgg	aggatgaata	tggcggattt	ctgaggccaa	ccattgtga		469

<210> 47
 <211> 349
 <212> DNA
 <213> Pinus radiata

<400> 47

ctgggtgtgat	ggcaggaatt	ccagtcctaa	ggccattttg	catctgtttg	ctttcagtct	60
acatgctgca	cattgtagct	gcagtagctt	caccaaggct	aggtagaagc	agcttcccaa	120
ggggtttcaa	atttggtgca	gggtcatctg	cttatcaggc	ggaaggagct	gctcatgagg	180
gtggcaagg	cccaagcatt	tgggatacat	tctccacac	tccaggtaaa	atcgctgatg	240
ggaagaatgg	ggatgttgca	gtagatcaat	accaccgtta	taaggaagat	gtgcagcttc	300

tcaaatacat ggggaatggac gtctatcggt tctctatctc ctgggtcacg

349

<210> 48
 <211> 385
 <212> DNA
 <213> Pinus radiata

<400> 48
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 aggaagaaag cgaaaatgtc aaagaaatta gcagagataa atttccggag agttttgaat 120
 tcggagttgc gacctccgcc tatcaagttg aagggtgctgc aaaaggagga ggaagaggcc 180
 ctagtatttg ggatacattt tcatatacac cagggaaaat tattgatgga agaaatgggtg 240
 atgttgcaat ggatcaatac catcgggtaca aggaggatgt ggatttaata gcaaaaatgg 300
 gattcaatgt gtatcgtttc tcaatatctt ggtctcgaat ctttccagat ggatttggag 360
 ctgaagtga taaggaagga atagc 385

<210> 49
 <211> 417
 <212> DNA
 <213> Pinus radiata

<220>
 <221> unsure
 <222> (209) ... (212)

<400> 49
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 tcatgttttg ggtagcaact tctgcttacc agtgtgaagg agctgcaaaa gaagggtggaa 180
 gaggtccaag catctgggac tcatTTTTnn nncagacacc aggcaaaatt gttgatggga 240
 gcaacgggtg tgtagctgtg gaccagtacc atcgttataa ggaagatgta aaacttataa 300
 aagatatggg agtggatgtc tacagattct caatctcatg gtctcgaatg tttccaaaag 360
 gaaaagggga gatcaatgag gaaggagtag cctattacaa taacctcata aatgaac 417

<210> 50
 <211> 264
 <212> DNA
 <213> Pinus radiata

<400> 50
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 agaagagttg cagacatgga tattaatggt cactgcaaga gctcctacaa atatagcagt 120
 gatcaagtac tgggggaaaa gagatgaaaa gctgatcctt cccatcaatg acagcatcag 180
 ctttactttg gatccagacc atctgtcagc cacaaccact gtagcagtta gccatcatt 240
 cacatctgat agaatgtggc tcaa 264

<210> 51
 <211> 417
 <212> DNA
 <213> Eucalyptus grandis

<400> 51
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 tttgcaatat ccctaaatga gcttacacca ggcttacaaag aaaaactgcc tccaactgat 120
 tcaaggttga gacctgatca gagacactta gaaaatgggg aatatgagtt ggcaaatgcc 180
 gagaagttaa gactggaaca catcacagaga caggcaagaa agttacagga gggaggttgg 240
 caaccgcatg gggttgggaa ggatgatgat ggatgttacc gctacatggg tgggtattgg 300
 gaagctcgag aagcatacga actgggatgg aatccctgac atattcgggc aaaaatgttg 360
 atgcttcaac ctggcactgt ggtaggatag atatgtcctc ctgcttgctt gaatata 417

<210> 52

<211> 305
 <212> DNA
 <213> Pinus radiata

<400> 52
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 ggagggatca cttttatcgc agagcaggtc agccaccatc ctccaatggg ctcagcctat 120
 gcagaaaaatg aacattttac atacagtctg tcctcaaaag taaaaaccaa gtttcttggc 180
 aactctgtgg atattttacc acttggaagg acacgtgtgg tgctaaagaa atccggagac 240
 gttctagatt tgggtgccgc tccatctaaa gttcataacc taatttttgg acgaacttgg 300
 attga 305

<210> 53
 <211> 474
 <212> DNA
 <213> Eucalyptus grandis

<400> 53
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 cggaagcaag cggccatggc gagcgactcg agcgcgaccc agctcaaate cgacgccttg 120
 atggagcaga tgaagcagca cctctccacc gagcgcggca aggcgggtcac caagaagatc 180
 ggcctcgtct accagatcaa catcgccccc aagaaaattg ggttcgacga ggtggtctac 240
 atcgtcgatc tgaagaagg agaggtcact aaaggaccat atgaagggtg aaaacctgat 300
 gctacctttt ctttcaaaga tgatgatttt atcaagggtg ccacaggaaa aatgaatcct 360
 caaattgctt ttatgagggg agcaatgaag attaaggga gcttgagtgc agcgcagaaa 420
 ttcactcctg acatattccc aaagccatcg aagatgtgag cattttgaaa agg 474

<210> 54
 <211> 562
 <212> DNA
 <213> Pinus radiata

<400> 54
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 acactagtgg atccaaagaa ttcggcacga gctttgaggc aacctacatt cattgaatcc 120
 caggatttct tcttgctcaa acaggtttta ggaaatggca ggcacaagtg ttgctgcagc 180
 agaggtgaag gctcagacaa cccaagcaga ggagccgggt aaggttgctc gccatcaaga 240
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 cgtgtaccct cgtgagcccg agccaatgaa ggagctccgc gaagtgactg ccaagcatcc 360
 ctggaacctc atgactactt ctgccgatga ggggtcaattt ctgggcctec tgctgaagct 420
 cattaacgcc aagaacacca tggagattgg ggtgtacact ggttactcgc ttctcagcac 480
 agcccttgca ttgcccgatg atggaaagat tctagccatg gacatcaaca gagagaacta 540
 tgatatcgga ttgcctataa tt 562

<210> 55
 <211> 1961
 <212> DNA
 <213> Pinus radiata

<400> 55
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 aatcaattga aagggtttta ttttcagtat ttcgatcgcc atggccaacg gaatcaagaa 120
 ggtcgagcat ctgtacagat cgaagcttec cgatatcgag atctccgacc atctgcctct 180
 tcattcgtat tgctttgaga gtagtagcga attcgcagac agaccctgtc tgatcgatgg 240
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 cggctctggc agctcgggt tgcagcagg gacaggtgtc atgcttctcc ttccgaattg 360
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 caatcctttc tacaagccgg gcgagatcgc caaacaggcc aaggccgcgg gcgcgcgcga 480
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 aagccgacga aaccatgc cggccgtga caatccacc ggacgatgtc gtggcggttc 660

cctattcttc	cgggaaccacg	gggctcccca	agggcgtgat	gttaacgcac	aaaggcctgg	720
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acgtgatact	ctgtgtcttg	cctcttttcc	acatctattc	tctcaattcg	gttctcctct	840
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gtgcagagta	agcgccctat	aaggagagag	agagcttata	aattgtatca	tatggattgt	1860
caacgcccta	cactcttgcg	atcgctttca	atatgcatat	tactataaac	gatatatgtt	1920
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<210> 56

<211> 414

<212> DNA

<213> Eucalyptus grandis

<400> 56

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cattgaactc	tctctctctc	tctctctctc	tctctctctc	tccccaccc	ccccttccca	120
accccaccca	catacagaca	agtagatagc	cgcacacaga	agaagaaaag	atgggggttt	180
caatgcagtc	aatcgacta	gcgacgggtc	tggccgtcct	aacgacatgg	gcgtggaggg	240
cgggtgaactg	ggtgtggctg	aggccgaaga	ggctcgagag	gcttctgaga	cagcaaggtc	300
tctccggcaa	gtcctacacc	ttcctggctg	gcgacctcaa	ggagaacctg	cggatgctca	360
aggaagccaa	gtccaagccc	atcgccgtct	ccgatgacat	caagcctcgt	ctct	414

<210> 57

<211> 469

<212> DNA

<213> Eucalyptus grandis

<400> 57

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ctcatctcct	agcagttcta	gggggttggt	tgtcctctgt	aattctatgg	agggcaagat	120
cttctccgaa	caaaccacaa	ggtactgcct	tacccccgga	gctgccgggc	gcatggccga	180
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gcccacaaatc	caaggcggga	atccacttgg	gctacgggta	tgccgggttt	ggcttcgtag	420
aatacgggga	cttttggcgc	gagatgagga	agatcaccat	gctcgagct		469

<210> 58

<211> 760

<212> DNA

<213> Eucalyptus grandis

<400> 58

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aaggagctta	agaagcatca	tcaatggcag	ccaacgcaga	gcctcagcag	acccaaccag	120
cgaagcattc	ggaagtcggc	cacaagagcc	tcttgacagag	cgatgctctc	taccagtata	180

tattggagac	cagcgtctac	ccaagagagc	cagagcccat	gaaggagctc	agggaaataa	240
cagccaaaca	tccatggaac	ctgatgacca	catcggcgga	tgaaggcag	ttcctgaaca	300
tgctcctcaa	gctcatcaac	gccaaagaac	ccatggagat	cggcgtctac	accggctact	360
ctctcctcgc	aaccgcccct	gctcttcccg	atgacggaaa	gatcttggcc	atggccatca	420
atagggagaa	cttcgagatc	gggctgcccc	tcatccagaa	ggccggccct	gcccacaaga	480
tcgatttcag	agaaggccct	gcccgtgccg	tccttgatca	gctcgtgcaa	gatgagaaga	540
accatggaac	gtacgacttc	ttctcaatcc	ttaatcggtc	atttgaatac	aaatacatgc	600
tcaatgggtc	aaagacaaca	taagacagaa	gatggaaaaa	atagaaagga	agggaaagtat	660
taagggtagt	ttctcatttc	atcaatgctt	gatcttgaga	tctcctttct	gggtcgatca	720
gctgaccg	cgccacaggt	gatgccatcc	ccgacgggaa			760

<210> 59

<211> 468

<212> DNA

<213> Eucalyptus grandis

<400> 59

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aaacaaatgg	gttccgcccg	attcgaatcg	gccacaaagc	cgcacgcctg	ttgcattccc	120
taccctgcac	aaagccacat	tggcgccatg	ctcaagctag	caaagctcct	ccatcacaaag	180
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aactatatgg	tcagcccat	caacgatctt	gtatcgagcc	tgggctcgaa	cccagagcgtc	420
cctccggtga	cttgcacaa	tctcggtatg	tttcatgaca	ctcgtgac		468

<210> 60

<211> 684

<212> DNA

<213> Pinus radiata

<400> 60

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ttgttggacg	ccatggaagc	tctccgaaa	gccgggatcc	tggaaaccgt	taaactgcag	180
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gacatggctc	tccacggcgg	caacttccag	ggaacaccca	tcggagtctc	catggacaac	660
atgcgaatct	ctttggcagc	cgtc				684

<210> 61

<211> 479

<212> DNA

<213> Pinus radiata

<400> 61

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cgagcaagga	agaaaaatag	gttgacgacg	cagaaattac	gcaggccaat	gaagttcaag	120
ttaaaagcac	tgggctgtgc	acggacttcg	gctcgtctgg	cagcgatcca	ctgaactggg	180
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gggtcgagga	gagttcaaac	tgggttctca	cccagatgac	caaggggacg	gatacctatg	420
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<210> 62

<211> 1785

<212> DNA

<213> Pinus radiata

<400> 62

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cacgaggttg	caggtcgggg	atgatttgaa	tcacagaaac	ctcagcgatt	ttgccaagaa	180
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tgaaccacca	ggacattcag	agcaaggtgc	gcgcagagct	ggacgctgtt	cttggaccag	1020
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aataaaagtt	tgcataaatt	aaatgatatt	tcaatatact	attttgactc	tccaccaatt	1740
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<210> 63

<211> 475

<212> DNA

<213> Pinus radiata

<400> 63

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gaaaatcttc	agtagcaacc	agcacaacat	cacttcggct	gaatacggcc	cgctgtggcg	420
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<210> 64

<211> 957

<212> DNA

<213> Pinus radiata

<400> 64

gaattcggca	cgagaaagcc	ctagaatttt	ttcagcatgc	tatcacagcc	ccagcgacaa	60
ctttaactgc	aataactgtg	gaagcgtaca	aaaagtttgt	cctagtttct	ctcattcaga	120
ctggtcaggt	tccagcattt	ccaaaataca	cacctgctgt	tgtccaaaga	aatttgaaat	180
cttgactca	gccctacatt	gatttagcaa	acaactacag	tagtgggaaa	atttctgtat	240

tggaagcttg	tgtcaacacg	aacacagaga	agttcaagaa	tgatagtaat	ttgggggttag	300
tcaagcaagt	tttgtcatct	ctttataaac	ggaatattca	gagattgaca	cagacatatc	360
tgaccctctc	tcttcaagac	atagcaagta	cggtagagtt	ggagactgct	aagcaggctg	420
aactccatgt	tctgcagatg	attcaagatg	gtgagatttt	tgcaaccata	aatcagaaaag	480
atgggatggg	gagcttcaat	gaggatcctg	aacagtacaa	aacatgtcag	atgactgaat	540
atatagatac	tgcaattcgg	agaatcatgg	cactatcaaa	gaagctcacc	acagtagatg	600
agcagatttc	gtgtgatcat	tcctacctga	gtaagggtgg	gagagagcgt	tcaagatttg	660
acatagatga	ttttgatact	gttccccaga	agttcacaaa	tatgtaacaa	atgatgtaaa	720
tcatcttcaa	gactcgctta	tattcattac	tttctatgtg	aattgatagt	ctgttaacaa	780
tagtactgtg	gctgagtcga	gaaaggatct	ctcggtatta	tcacttgaca	tgccatcaaa	840
aaaatctcaa	atttctcgat	gtctagtctt	gattttgatt	atgaatgcga	cttttagttg	900
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<210> 65

<211> 471

<212> DNA

<213> Pinus radiata

<400> 65

gaattcggca	cgagaaaacc	ttttcagacg	aatgttctga	tgctcggccc	cgccagaca	60
acagacatac	ttctcactgc	caatcaggct	acaggtagat	actacatggc	tgctcgagca	120
tattccaacg	ggcaaggagt	tcccttcgat	aacaccacta	ccactgccat	tttagaatac	180
gagggaaagt	ctaagacttc	aactccagtc	atgcctaata	ttccattcta	taacgacacc	240
aacagtgcga	ctagcttcgc	taatggctct	agaagcttgg	gctcacacga	ccaccagtc	300
ttcgcttcctc	agagtgtgga	ggagaatctg	ttctacacca	tcgggttggg	gttgatcaaa	360
tgctcggggc	agtcctgtgg	aggtccaacg	gatcaagatt	tgcaagcaag	atgaatacat	420
atcatttgtc	ccgcaaccac	ttcttccaat	ccttcaagct	cagcattttg	g	471

<210> 66

<211> 1010

<212> DNA

<213> Pinus radiata

<400> 66

gacaaacttg	gtcgtttgtt	taggttttgc	tgcaagtgaa	cactaatatg	gaaggccaga	60
ttgcagcatt	aagcaaagaa	gatgagttca	tttttcacag	cccttttcct	gcagtacctg	120
ttccagagaa	tataagtctt	ttccagtttg	ttctggaagg	tgctgagaaa	taccgtgata	180
aggtggccct	cgtggaggcc	tcacacaggga	aggagtacaa	ctatggtcag	gtgatttcgc	240
tcacaaggaa	tggtgcagct	gggtcgtgg	acaaaggcat	tcaaaagggc	gatgtgttat	300
ttgttctgct	tcctaatatg	gcagaatacc	ccattattgt	gctgggaata	atgttgcccg	360
gcgcagtgtt	ttctggggca	aatccttctg	cacacatcaa	tgaagttgaa	aaacatatcc	420
aggattctcg	agcaaagatt	gttgtgacag	ttgggtctgc	ttatgagaag	gtgaggcaag	480
tgaaactgcc	tgttattatt	gcagataacg	agcatgtcat	gaacacaatt	ccattgcagg	540
aaatttttga	gagaaactat	gaggccgcag	ggccttttgt	acaaatttgt	caggatgatc	600
tgtgtgcact	cccttattcc	tctggcacca	caggggcctc	taaagggtgc	atgctcactc	660
acagaaatct	gattgcaaat	ctgtgctcta	gcttggttga	tgtccatgaa	tctcttgtag	720
gaaatttcac	cacgttgagg	ctgatgccat	tctttcacat	atatggcatc	acgggcatct	780
gttgccgcac	tcttcgcaac	ggaggcaagg	tcgtgggtcat	gtccagattc	gatctccgac	840
actttatcag	ttctttgatt	acttatgagg	tcaacttcgc	gcctattgtc	ccgcctataa	900
tgtcttcctc	ccggtttaaa	aatcctatcg	ttaacgagtt	cgatctcagc	cgcttgaaac	960
tcctaaagctg	ttcatgactg	cggctgctcc	actggcgccg	gatctactgc		1010

<210> 67

<211> 1410

<212> DNA

<213> Pinus radiata

<400> 67

gaagatgggg	ctgtgggtgg	tgctggcttt	ggcgctcagt	gcgcactatt	gcagtctcag	60
gcttacaaatg	tggttaagtcc	aagcaatgct	actgggagtt	acagtgaagaa	tggtattggtg	120
atgaattact	atggggactc	ttgccctcag	gctgaagaga	tcattgtctga	acaagtacgc	180

ctgttggtaca	aaagacacaa	gaacactgca	ttctcatggc	ttagaaatat	tttccatgac	240
tgtgtgtgtg	agtcattgtga	tgcattcgctt	ctgttggact	caacaaggaa	cagcatatca	300
gaaaaggaca	ctgacaggag	cttcggcctc	cgcaacttta	ggtatttggg	taccatcaag	360
gaagccgtgg	agagggagtg	ccccggggtc	gtttcctgtg	cagatatact	cgttctctct	420
gccagagatg	gcgttgtatc	gttgggagga	ccatacattc	ccctgaagac	gggaagaaga	480
gatggacgga	agagcagagc	agatgtgggtg	gagaattacc	tgcccgatca	caatgagagc	540
atctccactg	ttctgtctcg	cttcaaagcc	atgggaatcg	acaccctggg	ggttgttgca	600
ctgctggggg	ctcacagcgt	ggggaggact	cactgcgtga	agctgggtgca	caggctgtac	660
ccggaagtag	atccgacact	ggaccctggg	cacgtggagc	acatgaagca	caagtgcccg	720
gacgcgatcc	ccaaccgaa	ggcagtgcag	tatgtgcgga	acgaccgggg	aacgcctatg	780
aagctggaca	acaactacta	cgtgaacctg	atgaacaaca	aggggctcct	aatagtggac	840
cagcaactgt	atgcagattc	gaggaccagg	ccgtatgtga	agaagatggc	aaaaagccag	900
gaatacttct	tcaaatactt	ctcccggggc	ctcaccatcc	tctctgagaa	caatcctctc	960
accggcgctc	gaggagaaat	ccgtcggcag	tgctcgctca	aaaacaaatt	gcacacaaaa	1020
agcaagcgtt	gagcgatagc	tcaatgccgc	agtgggtggg	gtgatagcgt	gatgccacag	1080
tgggtgggcat	ttcatatata	aattgcagtt	tgctgtttta	ttagataatc	ataatggtgt	1140
gggtgtgacta	tgccctgcga	atcacatcga	tgaaccacaa	ccgaaccgtg	gaacagttagg	1200
cttattccct	tatgtaagca	gaacctttta	ttataagcaa	aaaagacaat	cctgtctgtt	1260
attctagtat	aattttgtca	tcagttaaag	ttgctcatct	gataataact	ggaaacggta	1320
aaatatgaca	actacgtatc	ttctttggtc	atctgataat	aaccggaaac	gataaaaat	1380
gacaactaca	tatatctttt	aaaaaaaaaa				1410

<210> 68

<211> 607

<212> DNA

<213> Eucalyptus grandis

<400> 68

gaattcggca	cgagccaacc	ctggaccagg	tacttttggc	aggcggtcca	ttgcccttca	60
aaccggtcca	aaccggacca	tcactgtcct	tatatacgtt	gcatcatgcc	tgctcataga	120
acttaggtca	actgcaacat	ttcttgatca	caacatatta	caatattcct	aagcagagag	180
agagagagag	agagagagag	agagagagag	agagtttgaa	tcaatggcca	ccgccggaga	240
ggagagccag	acccaagccg	ggaggcacca	ggaggttggc	cacaagtctc	tccttcagag	300
tgatgtctct	taccaatata	ttttggagac	cagcgtgtac	ccaagagagc	ctgagcccat	360
gaaggagctc	agggaaataa	cagcaaaaaca	tccatggaac	ataatgacaa	catcagcaga	420
cgaagggcag	ttcttgaaca	tgcttctcaa	gctcatcaaa	gccaaagaaca	ccatggagat	480
tgggtgtcttc	actggctact	ctctcctcgc	caccgctctt	gctcttcctg	atgacggaaa	540
gattttggct	atggacatta	acagagagag	ctatgaactt	ggcctgccgg	catccaaaaa	600
gccggtg						607

<210> 69

<211> 421

<212> DNA

<213> Eucalyptus grandis

<400> 69

gaattcggca	cgagccgttt	tatttctctt	gatttctctt	gctcgagtct	cgcggaagag	60
agagaagaga	ggagaggaga	gaatgggttc	gaccggatcc	gagaccacga	tgaccccgac	120
ccaagtctcg	gacgaggagg	cgaacctctt	cgccatgcag	ctggcgagcg	cctccgtgct	180
ccccatggtc	ctcaaggccg	ccatcgagct	cgacctcctc	gagatcatgg	ccaaggccgg	240
gccggggcgc	ttcctctccc	cgggggaagt	cgcggcccag	ctcccgaacc	agaacccga	300
ggcaccgcga	atgctcgacc	ggatcttccg	gctgctggcc	agctactccg	tgctcacgtg	360
caccctccgc	gacctccccg	atggcaaggt	cgagcggtc	tacggcttag	cgccgggtgtg	420
c						421

<210> 70

<211> 508

<212> DNA

<213> Eucalyptus grandis

<400> 70

gaattcggta	cccgggttcg	aaatcgataa	gcttggatcc	aaagaattcg	gcacgagatc	60
actaaccatc	tgcctttctt	catcttcttt	cttctgcttc	tcctccgttt	cctcgtttcg	120
atatcgtgaa	aggagtccgt	cgacgacaat	ggccgagaag	agcaagggtcc	tgatcatcgg	180
agggacgggc	tacgtcggca	agttcatcgt	ggaagcgagt	gcaaaagcag	ggcatcccac	240
gttcgcgctg	gttaggcaga	gcacgggttc	cgaccccgtc	aagggccagc	tcgtcgagag	300
cttcaagaac	ttgggcgtca	ctctgctcat	cgggtgatctg	tacgatcatg	agagcttggt	360
gaaggcaatc	aagcaagccg	acgtgggtgat	atcgacagtg	gggcacatgc	aaatggcgga	420
tcagaccaa	gaatcgtcga	cgccattaaa	ggaagctggc	aacgttaagg	tttggtgggt	480
ggttcatttg	atctgggttg	ggggggtc				508

<210> 71

<211> 495

<212> DNA

<213> Eucalyptus grandis

<400> 71

gaattcggca	cgagggttaat	ggcagtgacg	cctcaacacc	acccaccttc	ctccatctct	60
ctcctccctt	cttctttctc	tgacttcaat	ggcagccgac	tccatgcttg	cgttcagtat	120
aagagggaag	tggggcagcc	taaaggggca	ctgcgggtca	ctgcatcaag	caataagaag	180
atcctcatca	tgggagcgac	ccgtttcatc	ggtgtgtttt	tgtcgagact	acttgtcaaa	240
gaaggtcatc	aggtcacttt	gtttaccaga	ggaaaagcac	ccatcactca	acaattgcct	300
ggtgagtcgg	acaaggactt	cgtgatctt	tcattccaaga	tcctgcattt	gaaaggagac	360
agaaaggatt	ttgattttgt	taaatctagt	cttgctgcag	aaggctttga	cgttggtttat	420
gacattaacg	gcgagaggcg	gatgaagtcg	caccaatttt	ggatgcctgc	caaaccttga	480
accagtcaac	tactg					495

<210> 72

<211> 472

<212> DNA

<213> Eucalyptus grandis

<400> 72

gaattcggca	cgagcataag	ctctcccgta	atcctcacat	cacatggcga	agagcaaggt	60
cctcgtcgtt	ggcggcactg	gctacctcgg	gcggagggttc	gtgagggcga	gcctggacca	120
gggccacccc	acgtacgtcc	tccagcgtcc	ggagaccggc	ctcgacattg	agaagctcca	180
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ccacttcggg	agccacaaca	tcctgatgca	gctcaagctc	gtggaggcta	tcaaagaagc	360
tggaatatgtc	aagcgttttt	tgccgtcaga	gttcgggaatg	gaccgggcc	tcattgggtca	420
tgcaattgag	ccgggaaggg	tcacgttcga	tgagaaatgg	aggtgagaaa	ag	472

<210> 73

<211> 380

<212> DNA

<213> Eucalyptus grandis

<400> 73

ggcaaacacg	cccgttttcg	ttttactaag	agaagatggt	gagcgttgtg	gctggtagag	60
tcgagagctt	gtcgagcagt	ggcattcagt	cgatcccgca	ggagtatgtg	aggccgaagg	120
aggagctcac	aagcattggc	gacatcttcg	aggaggagaa	gaagcatgag	ggccctcagg	180
tcccgaacct	cgacctcgag	gacatagcgt	ctaaagaccc	cgtgggtgag	gagagggtgcc	240
acgaggagct	cagggaaggct	gccaccgact	ggggcgctcat	gcacctcgtc	aacctatggga	300
tccccaacga	cctgattgag	cgtgtaaaga	aggctggcga	ggtgttcttc	aacctcccgga	360
tcgaggagaa	ggacaagcat					380

<210> 74

<211> 515

<212> DNA

<213> Eucalyptus grandis

<400> 74

ctctctctct	ctctctctct	gtgtgttcat	tctcgttgag	ctcgtggctg	cctcccccca	60
tggatccgca	caagtaccgt	ccatccagt	ctttcaacac	ttctttctgg	actacgaact	120
ctgggtgctcc	tgtctggaac	aataactctt	cggtgactgt	tggaagcaga	ggtccaattc	180
ttcttgagga	ttatcacctc	gtggagaaac	ttgccaaact	tgatagggag	aggattccag	240
agcgtgtggt	gcatgccaga	ggagccagt	caaagggtt	ctttgaggtc	actcatgaca	300
tttcccagct	tacctgtgct	gatttccttc	gggcaccagg	agttcaaaca	cccgtgattg	360
tccgtttctc	cactgtcatc	cacgaaaagg	gcagccctga	aaccctgagg	gaccctcgag	420
gttttctgt	gaagttctac	acaagagagg	gtaactttga	tctggtggga	aacaatttcc	480
ctgtcttctt	tgtccgtaat	gggataaatt	cccc			515

<210> 75

<211> 487

<212> DNA

<213> Eucalyptus grandis

<400> 75

gaattcggca	cgagctccca	cttctgtctc	gccaccatta	ctagcttcaa	agcccagatc	60
tcagtttctg	gctctcttcg	tcattctctg	ctcttgccat	ggatccgtac	aagtatcgcc	120
cgtccagcgc	ttacgattcc	agcttttga	caaccaacta	cgggtgctccc	gtctgggaaca	180
atgactcatc	gctgactgtt	ggaactagag	gtccgattct	cctggaggac	taccatctga	240
ttgagaaact	tgccaacttc	gagagagaga	ggattcctga	gcgggtggtc	catgcacggg	300
gagccagcgc	gaaagggttc	ttcgaggtca	cccacgacat	ctctcacttg	acctgtgctg	360
atttcctccg	ggctcctgga	gtccagacgc	ccgtaatcgt	ccgtttctcc	accgtcatcc	420
acgagcgcg	cagcccgaac	ctcagggacc	ctcgtgggtt	tgcagtgaag	ttctacacca	480
gagaggg						487

<210> 76

<211> 1474

<212> DNA

<213> Pinus radiata

<400> 76

gaattcggca	cgagaaaacg	tccatagctt	ccttgccaac	tgcaagcaat	acagtacaag	60
agccagacga	tcgaatcctg	tgaagtgggt	ctgaagtgat	gggaagcttg	gaatctgaaa	120
aaactgttac	aggatatgca	gctcgggact	ccagtggcca	cttgctcccct	tacacttaca	180
atctcagaaa	gaaaggacct	gaggatgtaa	ttgtaaagg	catttactgc	ggaatctgcc	240
actctgattt	agttcaaatg	cgtaatgaaa	tggacatgtc	tcattaccca	atggctccctg	300
ggcatgaagt	gtgggggatt	gtaacagaga	ttggcagcga	ggtgaagaaa	ttcaaagtgg	360
gagagcatgt	aggggttggt	tgcattgttg	ggctcctgtc	cagttgctgt	aattgcaatc	420
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gcacacctac	tcagggcgga	tttgcaagca	gtatgggtgt	tgatcagatg	tttgtgggtc	540
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taatggacac	cattccagt	gtcatcctc	tggaaccata	tcttgccctt	ctgaagacaa	900
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ttcagatgtt	tttttaactt	gtatatgtaa	agatcaattt	ctcgtgacag	taaataataa	1380
tccaatgtct	tctgccaat	taatatatgt	attcgtattt	ttatatgaaa	aaaaaaaaaa	1440
aaaaaaaaaa	aaaaaaaaaa	aaaaaaaaaa	aaaa			1474

<210> 77

<211> 414

<212> DNA

<213> Eucalyptus grandis

<400> 77

cacgctcgac	gaattcggta	ccccgggttc	gaaatcgata	agcttggtac	caaagcaaca	60
cattgaactc	tctctctctc	tctctctctc	tctctctctc	tccccaccc	ccccttccca	120
acccccacca	catacagaca	agtagatacg	cgcacacaga	agaagaaaag	atggggggtt	180
caatgcagtc	aatcgacta	gcgacggttc	tggccgtcct	aacgacatgg	gcgtggaggg	240
cgggtgaactg	ggtgtggctg	aggccgaaga	ggctcgagag	gcttctgaga	cagcaaggtc	300
tctccggcaa	gtcctacacc	ttcctggctg	gcgacctcaa	ggagaacctg	cggatgctca	360
aggaagccaa	gtccaagccc	atcgccgtct	ccgatgacat	caagcctcgt	ctct	414

<210> 78

<211> 273

<212> DNA

<213> Eucalyptus grandis

<400> 78

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catcttcccc	tacctgcccc	tcccagctca	ccatcgtcgc	gataaggctc	ggaagaagct	120
ttctgagatt	tttgcaaaca	tcattttctc	acgaaaatgt	gctggcaa	cagaagaaga	180
catgttgag	tgcttcattg	actccaagta	caaaaatgg	cgcccgacaa	ctgaggccga	240
ggctactgg	ctgcttattg	cggtctctct	tgc			273

<210> 79

<211> 121

<212> DNA

<213> Eucalyptus grandis

<400> 79

ctacctctc	accaacaaga	agtacctctc	tgccgtctct	aatgaacaga	agcacctgat	60
ggagaagcat	gggaacaatg	ttgatcatga	tggtctttct	gaaatggatg	tcctgtatcg	120
g						121

<210> 80

<211> 505

<212> DNA

<213> Eucalyptus grandis

<400> 80

ggaggctgaa	gatagcagaa	ggagaggacg	gtccctacct	gtacagcacc	aacaactacg	60
tggggagaca	gatttgggag	ttcgatccgg	aggctggcac	cgctgaggag	cgcgcgagg	120
tcgaggctgc	tcgccagcac	ttctacgacc	accgccacca	agtcaagccc	tgtggcgacc	180
tcctctggcg	catgcagttc	ctgagggaga	aggagttaa	gcagacgatt	ccgccagtga	240
gggtggagga	tggcgaggag	atcacctacg	acaaggcgctc	caccgcgctg	aagcgggccc	300
tccatttctt	ctccgccttg	caggctagcg	acggctcattg	gcctgccgag	aacgccggcc	360
ctctcttctt	cctccctccc	ctggatcatg	gcgtctacat	caccggccac	cttgacgccg	420
tcttccccgc	cgagcatcgc	aaagagattc	tccgctacat	ctacaaccac	cagaatgaag	480
atggtggatg	gggcttgac	atcga				505

<210> 81

<211> 270

<212> DNA

<213> Eucalyptus grandis

<400> 81

catggatgac	attgtctctc	acgagtttga	gcaaaagagg	ggccatgtag	tatctgcagt	60
ggagttgctc	ataaaatata	gtggtgtctc	ggagcaggaa	gctgtggagg	aactccagaa	120
acgagtcatt	gatgcatgga	aggacaccaa	tgaagagttt	ctccgtccaa	ttgcgggtccc	180
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tggggataat	tacaccatt	ctgaaaccaa				270

<210> 82
 <211> 441
 <212> DNA
 <213> Eucalyptus grandis

<400> 82
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 ccatggagga tgatcgagat cgggggcttc tttacgattc agatccgccg tctccatcgc 120
 tctccccgcc actctccccg ccgcggccgt tcgcgctcac cttcttcgac agggagaggc 180
 acgtgacgtt cctggagatg atgtaccaca tgctccctcg cccctaccag tcgcaggaga 240
 tcaaccacct caccctcgcc tacttcgtca tctccggcct cgacatcctc gacgccctcg 300
 atcgcgtaga caaagatgcg gttgctgact ggggtttatc tttccaagct catccgagga 360
 gtaaagctga tctagacaat ggacaatttt atgggtttca tggttccaga agctcacagt 420
 tcccttcaaa agataatgca a 441

<210> 83
 <211> 467
 <212> DNA
 <213> Eucalyptus grandis

<400> 83
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 aattttatgg gtttcatggt tccagaagct cacagttccc ttcaaaa 467

<210> 84
 <211> 396
 <212> DNA
 <213> Pinus radiata

<400> 84
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 atggaatatg gttggcatat aaatttgcca agattggaag caaggaaact catcgacgtg 180
 tttggacagg accccattta tttgatgcca aatatgaaga cccaaaaact tctagaactc 240
 gcaaagtggg agttcaatat gtttcaactt ttacaacagc aagagctaaa acttctctcc 300
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 tattacactt tggcatcttg cattgatagt gaacct 396

<210> 85
 <211> 462
 <212> DNA
 <213> Pinus radiata

<400> 85
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 caacagaaag agttaagca tgtgtccaga tgggtgaaag at 462

<210> 86
 <211> 247

<212> DNA

<213> Pinus radiata

<400> 86

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caaatattct	gcacatcata	tttaaaggaa	gccctaaaaa	cggttccgat	ctccaatggg	120
agtctttcag	gagagattga	atacgttatt	gaatatgggt	ggctcacaaa	tttcccgaga	180
ttggaagcac	gaaattatat	cgacgtattt	ggaaaggaca	ccattccctg	tgtaagacg	240
acgacca						247

<210> 87

<211> 426

<212> DNA

<213> Pinus radiata

<400> 87

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tgcaaaattg	gagttcaata	tctttcactc	cttacagcaa	aaagagttaa	aacagctgtc	180
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ctttgccaaa	acgtgttata	ttggaatagt	tctggacgac	atctatgaca	ctttcggaac	360
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gttcct						426

<210> 88

<211> 488

<212> DNA

<213> Pinus radiata

<400> 88

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aaattggagt	tcaatatctt	tacttcctta	caacaaaaag	agttaaaaca	gctctccaga	180
tggtggaaag	attcgggttt	ctctcgactc	acattcactc	ggcatcgta	cgtggaattc	240
tacactttgg	cctcctgcat	tgccactgag	cccaaacatt	cagcatttag	attgggcttt	300
gccaaaacgt	gttatcttgg	aatagttctg	gacgacatct	atgacacttt	cggaacgatg	360
gaggagctcg	aactcttcac	agccgcaatt	aagagatggg	atccttccgc	cagggagtg	420
cttcagaat	atatgaaagg	catatatatg	gtgttttacg	atgcgttaat	caaatggctc	480
gagaggcg						488

<210> 89

<211> 223

<212> DNA

<213> Pinus radiata

<400> 89

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cacatactct	gccaaataga	ttatcgatcc	aaatttgc	atctcatagg	tttgataggg	180
cgtttgctga	atgataccaa	aacttaccag	gcggagcgag	gtc		223

<210> 90

<211> 318

<212> DNA

<213> Pinus radiata

<400> 90

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cggaaaattg	cagaagactg	gtttttgaca	atgcaagatc	atcgcaactg	ttttgcatgg	120
aaaaatgacg	tttcacacat	tcacatgaaa	cggaaattaa	agagcatgtc	aaaaagatac	180

tggttcgaacc	agttgcgtag	aatacgttac	tcaaataaag	gcccggcact	ttcattttgt	240
actggttacca	cacatacttg	ctcagagggt	attatgaccg	ttctgttatt	cgtattttct	300
aatgcccgcga	atgaaatt					318

<210> 91
 <211> 1695
 <212> DNA
 <213> Eucalyptus grandis

<400> 91

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atcaagtcgg	tcgtcgaggg	aaagatgccg	tcgtactcgc	tcgagtccaa	gcttggggac	180
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ccggttgggt	atgtgcagat	accggtgggg	attgccgggc	cgctgttgct	cgacgggagg	360
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cgtccttaaa	gttta					1695

<210> 92
 <211> 315
 <212> DNA
 <213> Eucalyptus grandis

<400> 92

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ctgaacctgc	taggggtgaa	aggcgaaaag	gagctagcgg	gagccaactc	gaggctcctg	120
gccacggtcg	tgccggcgcg	cgtccttgcc	gccgagctct	ccctcatgtc	cgctatcgcg	180
gcggggcagc	tcgtgaagag	ccacatgaag	tacaacagggt	ccagcaaaga	cgttaccaag	240
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ggccacggaa	gaatc					315

<210> 93
 <211> 244
 <212> DNA
 <213> Eucalyptus grandis

<400> 93

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ggatctccgg	aaacttctgc	tcggataaga	agcccgacgc	tgtgaactgg	attgagggga	180
ggggcaagtc	ggttgtgtgc	gaggccgtga	tcaagggcga	cgtggtgagg	aagggtgctca	240
agac						244

<210> 94
 <211> 244
 <212> DNA
 <213> Eucalyptus grandis

<400> 94						
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gaaggacttc	cccgcacatg	acgtgatggg	gatctccgga	aacttctgct	cggataagaa	180
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caag						244

<210> 95
 <211> 419
 <212> DNA
 <213> Pinus radiata

<400> 95						
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gccggtgctc	ttggtggttt	taatgcccat	gcaagcaaca	tagtctctgc	aatatatata	180
gcaactggac	aggatccagc	tcagaatggt	gagagttctc	attgcatcac	catgatggaa	240
gctgttaatg	aggggaaggga	tcttcacata	tctgtcacca	tgccttccat	agaggttggc	300
actgttggag	gtggtactca	gcttgcgtct	cagtctgctt	gcctgaacat	gcttggcgctc	360
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<210> 96
 <211> 856
 <212> DNA
 <213> Eucalyptus grandis

<400> 96						
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<210> 97
 <211> 863
 <212> DNA
 <213> Eucalyptus grandis

<400> 97						
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<210> 98

<211> 668

<212> DNA

<213> *Eucalyptus grandis*

<400> 98

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<210> 99

<211> 430

<212> DNA

<213> *Pinus radiata*

<400> 99

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aagatgggga	atcaatcact	cccctcgatg	atctgattca	aggccttttg	atggtcgaca	420
gtgttgaacg						430

<210> 100

<211> 478

<212> DNA

<213> *Pinus radiata*

<400> 100

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<210> 101
 <211> 204
 <212> DNA
 <213> Pinus radiata

<400> 101
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 gtgatacagt ttctgtcaac gccttatggg gaactcgctt accgtgaacg tgctgagcga 180
 ctgattgatg aagtaaggaa cata 204

<210> 102
 <211> 299
 <212> DNA
 <213> Pinus radiata

<400> 102
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 gattgaggaa gtaaagaagg tattcaattc aatgtcagag gagaatggcg aattaatcac 180
 tccctgaat gatctgattc aacgcctttg gatggctgac agtgttgaac gtttggggat 240
 cgatagacat ttcgaaaatg agattgaatc agcgtggat tatgtttaca gttattgga 299

<210> 103
 <211> 399
 <212> DNA
 <213> Pinus radiata

<400> 103
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 gcttatcatt tataagttat gttcagacaa catataagt 399

<210> 104
 <211> 672
 <212> DNA
 <213> Pinus radiata

<400> 104
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 tgtgacagtt gcagatgcca gattctgcaa gccacttgat cgtgatctaa ttcgatctct 600
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 acatgttgct ca 672

<210> 105
 <211> 971
 <212> DNA
 <213> Pinus radiata

<400> 105

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attatccgat	tcatatgaag	aatctctcta	ttcgagagct	caaacaactt	tcaaataaac	300
ttcgttctga	cataatTTTT	gaggtttcaa	gaaccggtgg	ccaccttggg	tctagccttg	360
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tatgggacgt	tggccatcag	gcttatcctc	acaagattct	tactggtaga	agagataaga	480
tgcccacatt	gagacagaca	aatggcctct	caggctttac	aaaacgttca	gagagtgaat	540
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ttggtcgtga	cttgaaaggg	gaaaacaatc	atgtcattag	tgctcattga	gatggtgcca	660
tgacagctgg	gcaagccttt	gaagctatga	acaatgcagg	atatttggat	tccaacatga	720
tcgttattct	gaatgacaac	aaacaagttt	ctctgccaac	agcaaactct	gatgggccta	780
taccaccagt	gggtgcactc	agcagtgcac	tgagtaagct	tcaatcaagt	aaacctttgc	840
gtgaactaag	agagggttgc	aagggtgtta	ccaaacaact	cgggtccccc	atgcatgaac	900
tggcagcaaa	agtggacgag	tatgcacgag	gtatgatcag	tggttcccgt	tccacgctgt	960
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<210> 106

<211> 265

<212> DNA

<213> Eucalyptus grandis

<400> 106

aggcgggtcat	ctgagcgcca	gcctcggcgt	gggtggaactc	acgggtcgctc	ttcacaatgt	60
tttcaatgcc	ccagaagaca	aaatcgtatg	ggacgtggga	catcagacat	acccccacaa	120
gattctgacc	ggacggcgga	ctcggatgca	cacgattagg	aagacctcgg	ggcttgccggg	180
gtttcccaag	agggatgaga	gcgtttatga	tacatttggc	gtgggacaca	gttctacgag	240
catctccgcc	ggctcggta	tggcg				265

<210> 107

<211> 295

<212> DNA

<213> Eucalyptus grandis

<400> 107

ccccgtccg	ggaaaagctt	gtcaaagcat	ggagaaatga	ctctgagatc	tttgctcact	60
atggccggct	caccacgcca	tactctgatg	agcttctcgg	gagcaaattt	tgcttccatg	120
tcaaaggctt	tgaagtaaac	acagcccgca	ttgcagattc	attgtattat	ggctgtgtcc	180
ctgtgataat	cgccaatcac	tatgatctcc	cattcgcgga	catattgaac	tggaagagct	240
tctcggctcgt	ggttgccact	ttggacattc	cattgcttaa	gagaatcctc	aaggg	295

<210> 108

<211> 456

<212> DNA

<213> Eucalyptus grandis

<400> 108

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ctttttgatg	ctatagttag	cttgtgcac	ccagttatcg	tcagtgcacg	tatcgagttg	120
ccatttgaag	atgtcataga	ctaccggaaa	attgcaatat	ttgtagatac	tgccacttct	180
ctgaagcgag	gatttttggg	gaaacttctg	agaaaagtac	ggacagaaaa	gattctggaa	240
tatcagaaa	agctaaaaga	ggtgaagcga	tttttcgagt	atgggtgatcc	aaacgggaac	300
gtcaaggaaa	tttggcgcca	aatatcgag	aagctacccc	ttattaaaact	gatgattaac	360
cgtgacaaga	ggattgtcaa	gagggacatg	agtgaaccag	actggtcctg	tatctgtctg	420
aaccagacgg	gggtcatttc	cactctatga	cacgag			456

<210> 109

<211> 640

<212> DNA

<213> Eucalyptus grandis

<400> 109

cctctctctt	ccgtttgtgc	attgggctca	cgtaagaaga	aagagagaat	atgtcgcagg	60
tctcagcaac	tccatgcgct	ccccgaata	aagaacagg	ccatgtgatc	gaacgtcggg	120
ccgcgggtta	tcacccagc	gtatggggg	actacttct	taaatatgat	tctccctcca	180
actcagtga	gttcaaattc	ctcggaagag	tggagggaca	aattgaggaa	ctgaaaggag	240
aggtgaagaa	gatgctgatt	gatgtcgtgg	acaagccctt	accaaagctt	cacttgattg	300
atcaaatacca	acgcttgagg	attgagtacc	atgttgaacg	tgaagtagat	gagcaattgg	360
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acacagtcgc	tctcatcttt	agattgctgc	gacaacatgg	ttacaatatt	tcatcagaga	480
tctttgacaa	attcaaagat	agcaacggga	acttccgaga	gtcgtcata	agtgatgtgc	540
ggggattgct	gagcctatat	gaagcttgcc	atttaagggtg	tcatggcgat	tcaatcttgg	600
acgaagcact	tccatttgct	acaactcacc	ttgaatcatc			640

<210> 110

<211> 396

<212> DNA

<213> Eucalyptus grandis

<400> 110

tctctcttcc	gtttgtgcat	tgggcacacg	taagaaaaaa	gagagaatat	gtcgcaggtc	60
tcagcaactc	catgcgctcc	ttcgaataaa	ggaacaggcc	atgtgatcga	acgccggctg	120
gcgggttatc	acccagcgt	atggggggat	tatttcctta	aatatgattc	tccctccaac	180
tcagtgaagt	tcaaattcct	cggaagagtg	gagggacaaa	ttgaggaaact	aaaaggagag	240
gtgaagaaga	tgctgactga	tatcatggat	aagcccttac	aaaagcttca	cttgatcgat	300
caaatccaac	gcttggggaat	tgagtaccat	tttgaacgtg	aaatagatga	gcaattagaa	360
caaatccaca	aaagttactc	tcgacttgat	catgaa			396

<210> 111

<211> 348

<212> DNA

<213> Eucalyptus grandis

<400> 111

gttcttcagc	tacctagatc	tagaaagtag	caaaaaaaga	aatgtctctt	ccgatttcaa	60
gagtcaccac	ttcttctcct	gctgaaaaaa	caagcctggt	ccctgaaggc	ggatcggcaa	120
tttttcatcc	aaccatattg	gcggattact	ttctcaaaca	tgcttccaac	tccaactcga	180
cgagttctga	tggcgtagta	gaggaacata	ttgagagatt	gaaaggagaa	gtgaggaaga	240
tggtgatggg	tgctatggat	aagccatcgc	aaaagttgaa	cttgattgat	cagatccaac	300
gcttgggatt	tgcttaccat	tttgaacatg	agatagatga	gcagctag		348

<210> 112

<211> 508

<212> DNA

<213> Eucalyptus grandis

<400> 112

cacaagcttc	ctccctcct	caattcacca	taaccagcca	tctctcttat	tctttaggca	60
cctctgctcc	tctctctccg	ccgccacctc	ctccacctcc	tccggtgctc	aattcgtgac	120
atgcactttg	aagattgaag	ctcaagagat	cgggagacgg	tcggcggaatt	ggcaacctaa	180
cgtttttgac	tacgactttc	tgagtcact	caatgttgat	tacacggagg	ataaatactc	240
agaggaagcc	caaaggttga	agaaagaagt	caaggttcta	ttcgacaaga	agatgaattc	300
ggtggccaag	ctcgagttca	ttgacgtggg	tcaaagacta	ggactaggat	accaatttga	360
gacggagatc	aagaacgctc	taagttccat	ctataacaac	gccgaagatg	ctcaactttt	420
ggatgatctc	tatgccgttt	cccttcgatt	ccggctactt	agacaacatg	gattcaacat	480
atcgcaagat	gcgtttcaaa	ggttttatg				508

<210> 113

<211> 398

<212> DNA

<213> *Eucalyptus grandis*

<400> 113

cctcgatccg	ccctaaccac	ccctctctct	cactcttcag	tcgccctcgc	tcctccttct	60
cctccccctc	tgccgtttcc	tcaggcactc	ggtttgcgaa	atgtgctttg	actattgaag	120
atgaagatac	cgcgagacgt	tcggcggaatt	ggaagcctag	cgtttgggac	tatggctttg	180
tcgagtcact	caatactgat	ttcccgggtg	ataaatatac	agagcaagtt	caaagggtga	240
aggaagaagt	caagggtcta	ttccacaggg	agatgaatca	ggtggccaag	ctcgagttca	300
ttgacgtggt	tcaaagatta	ggactaggat	accattttga	gacggagatc	aataactctc	360
tcagttccat	ctataacaac	actgaagatg	ttcaactt			398

<210> 114

<211> 432

<212> DNA

<213> *Pinus radiata*

<400> 114

cggaagccaa	ggcaatcaat	caaaaatttg	tttgccatta	caactctgca	tttactagtc	60
ttagccactt	cgaagaattt	caaatggcta	gcgtttctgt	taaagcggga	gcaacttcta	120
ctgtgtcttg	tggtttggcc	agcaacaact	tgatccgaag	gacggccaat	cctcatecca	180
atgtctggga	ttatgacttt	gtacattctc	ttaagtcgcc	ttacaatgat	tctagttaca	240
cagaacgtgc	ggagacactt	attggccagc	ttaaggatgat	gcttagtgct	gcgattggag	300
gtggagaatc	aatgattact	ccatctgctt	atgacacagc	atgggtagcc	aggggtgcctt	360
ccattgatgg	ctctgcttgc	cctcaatttc	cccagacagt	tgaatggatt	ttgaaaaatc	420
aattaaaaga	tg					432

<210> 115

<211> 363

<212> DNA

<213> *Pinus radiata*

<400> 115

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tcatactca	taccatggc	ctccgtcgtc	gaccaagccg	agctgtgcag	caaactctgt	120
agcatgagtt	cacctggtgt	acaaagacgc	acaggcgatt	accattccaa	tctctgggac	180
gacgagttca	tccagtcctt	ctcaacgcct	tatggggcac	cttcttaccg	cgaacgtgct	240
gatagacttg	ttgggggaagt	aaaggagatg	ttcaattcac	ttacagtact	cactccccac	300
aatgatctcc	ttgagcaact	ttggatggtg	gatagcgttg	aacgtttggg	aatcgatagg	360
cat						363

<210> 116

<211> 779

<212> DNA

<213> *Pinus radiata*

<400> 116

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accaagccga	gctgtgtagc	aaatctgtga	gcatgagttc	acctggtgta	caaagacgca	120
caggcgatta	ccattccaat	ctctgggacg	acgatttcat	ccagtcactc	tcaacgcctt	180
atggggcacc	ttcttaccgc	gaacgtgctg	atagacttgt	tggggaagta	aaagagatgt	240
tcaattcact	tacactactc	actccccca	atgatctcct	tcaacgcctt	tggtatgggtg	300
ataccgttga	acgtctggag	atcgataggc	atttcagaaa	tgaaataaaa	tcagcgttgg	360
actacgttta	cagctattgg	agcgaaaaag	gcattggatg	tggttagagag	agtgttgtta	420
ctgatctcaa	ctcaactgcc	ttgggggttc	gaactcttcg	actacacgga	tttccgggtg	480
cctcagatgt	tttgggaagt	ttcaaagatc	aaaatgggaa	gtttgcaggc	tgctctgcca	540
atgcagagac	agaggcagag	atgagagaca	ttctcaattt	atttagggcc	tcccttgttg	600
cctttctctg	ggagaaagtc	atggaaagag	ctcaaacatt	ctgtacgtca	tatttacaag	660
aagccctaaa	aactgttccg	atctccaatg	atagtctttc	acgagagatt	gaatacgtta	720
ttgaatatgg	ttggctcaca	aatttttccg	agattggaag	caaggaatta	catcgacgt	779

<210> 117
 <211> 1173
 <212> DNA
 <213> Pinus radiata

<400> 117
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 aatctctctt gagatgggtg aaagattatg gattccctga aataacattc tcacggcatc 180
 gtcacgtgga atactacact ttagcagctt gcattgcaaa tgatcctaaa cattctgcgt 240
 ttcgactagg atttggtaaa ataagtcata tgatcacgat tctcgacgat atctacgaca 300
 ccttcggaac aatggaggag ctggaactct taaccgcagc gtttaagaga tgggatccgt 360
 cttcgataga gtgtcttcca gattatatga aaggagtgt catggcgggt tacgacaaca 420
 tcaacgaaat ggcacgagag gcgcagaaaa ttcaaggctg ggatacagtc agctatgctc 480
 gaaaatcttg ggaggctttt attggtgctt atatacaaga agccaagtgg atttccagtg 540
 gttatcttcc cacgttcgac gagtacctcg agaatgggaa ggtagcttc ggctctcgca 600
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 aaattgactt tccaccgaaa ttcaatgatt tgatatgtgc catccttcga ctgaaagggtg 720
 aactcaatg ctacaaggct gacagggcgc gtggagaaga agcttcggcc gtatcgtgtt 780
 atatgaaaga caatcctgga ataacagagg aagatgctgt caatcaagtc aatgctatgg 840
 tcgataaactt aaccaaggaa ctgaattggg agttacttag acccgacagc ggtgttccca 900
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 aaactgtgcc tttgtagcca cacatgaaat gtacaataag ttcttaagtt tctgacttcg 1080
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 ccatatcatt atatggagga ggtggaggag ttg 1173

<210> 118
 <211> 1634
 <212> DNA
 <213> Pinus radiata

<400> 118
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 catggatata ctgtgtcttc ggatgcgttt gaacacttta aagaccaaatt gggacagttt 120
 agcgtcttcg ccaatgatac agagttagcag ataagaagcg tttttaattt atttcgagct 180
 tctctcattg cttttcccga ggaaaaagtt ctggaagagg ctgaaaaatt cgctgctgca 240
 tatttaaaag cagccctaca aacccttcca gtctcgggtc tttcacgaga aatacaatac 300
 gttttcgatt atcgttggca ctcaaatctg cctagactgg aagctaggag ttacgtcgac 360
 atccttgtag ataatacgat cagtggaaag ccagatgcga aactaaaaa acttttagaa 420
 cttgcgaaat tggagttcaa tttttccat tctctacaac agaaagagtt acaatgtctg 480
 tggagatggg ggaaagaatg ggggtgcccc gaactaacct tcgttcgaca tcgttacgtg 540
 gaattctaca ctttgggtctc tggcactgac atggtgcctg aacatgctgc attcagactg 600
 agctttgtta aaacgtgtca tcttatcacg attctggatg atatgtacga cacttcgga 660
 acaattgacg agctccgact cttcacagcc gcggttaaga gatgggatcc gtcggcgacg 720
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 ccaacgtttc aggagtactt cgagaacggg aaactcagtt ctgggtcatcg catagcgacg 960
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 aagaaaaatg cttttaacat tagcagaggt ttacatcact tctacaacta ccgagatggc 1320
 tacagtgttg ccagcaacga aactaaagat ctgggtgatca aaaccgttct tgaacctgtg 1380
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 ctttaggtta tgtgttttat tattgtactt ttgactctt ttggctcgat aggcctagggt 1500
 tgcaaaagggt tttcaggacc ttgcgcatta atatagttgt caataatatg taagttatga 1560
 attctcagtg aatacggttt gttatcagtt ttttggggaa tctaatttct attaaagaaa 1620
 aaaaaaaaaa aaaa 1634

<210> 119
 <211> 301
 <212> DNA
 <213> Eucalyptus grandis

<400> 119
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 agaagattgc ccaggagacc taaatttctg gtatcgccac atctcaaaag gggcttggcc 180
 tttttcgaca gcagatcacg gatggcccat ttcagactgc acagcagaag gattaaaagc 240
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 g 301

<210> 120
 <211> 433
 <212> DNA
 <213> Eucalyptus grandis

<400> 120
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 aacaaggagc tctgagcgcg gccaccatc tcccggttt cgcgtgggctt gcggcgagag 120
 tctctctggg gttcctgggg cgtcgtttct tgatcgtggg atcaggatgt ggaaactgaa 180
 agttgcggaa ggagcaaatc cttggctaag aagtctgaac aatcatgttg gtagacaaat 240
 ttgggagttc gatccaaatt gtggatcccc agaagagatt caggagattg aagaggctcg 300
 tgcaaacctc ttaaagcatc ggtttgagaa gaagcacagc tcagatttga tgatgcgaat 360
 tcagttttcc aaggaaaaata caggcagagt agttttacca ccagtaaaagg tgaaagactt 420
 ggatgaaatc aca 433

<210> 121
 <211> 596
 <212> DNA
 <213> Eucalyptus grandis

<400> 121
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 cactggccag gggactatgg cggacatag tttctcatgc ctggcttggg aattgccctt 120
 tctattactg gagcactaaa tgccgtcttg tctgagcaac ataaacaaga gatgtgccga 180
 tatctgtaca atcatcaaaa caaagatggc ggttgggggt tgacattga aggtccaagc 240
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 gatggacaag gggctatgga gaaggcacga aaatggattc tggaccatgg cagtgtctact 360
 gcaataacat catggggaaa aatgtggctt tcagtacttg gagcatttga gtggctcaggc 420
 aataacccat tgccccctga gatatggctt cttccttaca tgcttccgat tcatccagga 480
 agaatgtggg gccactgccg gatggtttat ttgccgatgt catacttgta tgggaagagg 540
 tttgtaagtc ccataacacc aaccgttttt gtccttgaaa aaaggaactt tatgca 596

<210> 122
 <211> 332
 <212> DNA
 <213> Eucalyptus grandis

<400> 122
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 tggcagataa atatacaga gagagagaga aagagagaat gtggaagctg aagatagcgg 120
 agggcgggccc gtggctgacc agcgtgaaca accacgtcgg gcggcagcat tgggagtttg 180
 accctgacgc cgggacaccg gaggagaggg cggaggtgga gagggtccga gatgagttca 240
 cgaggaaccg gttccgaatc aagcagagcg ctgatctttt gatgaggatg cagctcacia 300
 aggagaaccc aagcgggccc attcaccgcc gg 332

<210> 123
 <211> 293

<212> DNA

<213> Eucalyptus grandis

<400> 123

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tccttaagaa	aggacatgaa	tttctgaagg	aatctcagat	cgaccgaaac	ccctctggcg	180
acttaaagaa	aatgtaccgt	cacattttcca	aaggagcatg	ggctttctcg	gacaaagatc	240
atggatggca	agtttcggat	tgcacagcag	aaagtatgaa	gtgttgcccta	gtt	293

<210> 124

<211> 604

<212> DNA

<213> Eucalyptus grandis

<400> 124

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catctctctg	cctctctctc	tctctctctc	actccagcgc	gctgactggg	catttcggat	120
ccgtaccaga	tggacacgga	caacaaactc	ttcaatgtgg	gcgtcttgct	cgtggccact	180
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tccgatttga	gccagcaaga	agtgtaccaa	ttcaatgtgc	cgactttcgg	acctggagtt	480
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aggattaata	agctcaaggg	gtatgtcaat	cagatgggta	tggaagccga	ggactacttc	600
tcaa						604

<210> 125

<211> 515

<212> DNA

<213> Eucalyptus grandis

<400> 125

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tctctgcctc	tctctctctc	tcactccagc	gcgctgactg	ggcatttcgg	atccgtacca	120
gatggacacg	gacaacaaac	tcttcaatgt	gggcgtcttg	ctcgtggcca	ctctcgttgt	180
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taggacatgg	ccggtggttg	gtgggctgct	ccggttcttg	aaggggtccga	tgggtgatgct	300
gcgggaagag	taccccaagc	ttgggagcgt	attcactctg	aatctgttga	acaagaaaat	360
aacgttcttc	atcggccctg	aggtttctgc	gcacttcttc	aaggcttccg	agtccgattt	420
gagccagcaa	gaagtgtacc	aattcaatgt	gccgactttc	ggacctggag	ttgtattcga	480
cgtcgattac	accatcaggc	aagagcagtt	tcggt			515

<210> 126

<211> 366

<212> DNA

<213> Eucalyptus grandis

<400> 126

gctgactggg	catttcggat	ccgtaccaga	tggacacgga	caacaaactc	ttcaatgtgg	60
gcgtcttgct	cgtggccact	ctcgttggtg	ccaagcta	ctcggcgtct	attccgagat	120
ccggaagcgc	cctccctccc	gtcgttagga	catggccggt	ggttggtggg	ctgctccggt	180
tcttgaaggg	tccgatgggtg	atgctgcggg	aagagtaccc	caagcttggg	agcgtattca	240
ctctgaatct	ggtgaacaag	aaaataacgt	tcttcatcgg	ccctgaggtt	tctgcgact	300
tcttcaaggc	ttctgagtcc	gatttgagcc	agcaagaagt	gtaccaattc	aatgtgccga	360
cttttcg						366

<210> 127

<211> 458

<212> DNA

<213> Eucalyptus grandis

<400> 127

ttcttgaagg	gtccgatgg	gatgctgcgg	gaagagtacc	ccaagcttgg	gagcgtattc	60
actctgaatc	tgttgaaca	gaaaataacg	ttcttcatcg	gccctgaggt	ttctgcgcac	120
ttcttcaagg	cttccgagtc	cgatttgagc	cagcaagaag	tgtaccaatt	caatgtgccg	180
actttcggac	ctggagtgt	attcgacgtc	gattacacca	tcaggcaaga	gcagtttcgg	240
ttttttactg	aggctctgag	gattaataag	ctcaaggggt	atgtcaatca	gatggttatg	300
gaagcggagg	actacttctc	aaaatgggga	gatagtggcg	agggtggacct	aaagtatgag	360
cttgagcaact	tgaccatatt	gacagcgagc	agatgtcttt	tgggtcgaga	ggttcgtgag	420
aagctctttg	atgatgtgtc	agccctcttc	cacgacct			458

<210> 128

<211> 442

<212> DNA

<213> Eucalyptus grandis

<400> 128

ctttgatgat	gtgtcagccc	tcttccacga	ccttgacaat	ggaatgctac	cgatcagtgt	60
catcttcccc	tacctgcccc	tcccagctca	ccatcgctcg	gataaggctc	ggaagaagct	120
ttctgagatt	tttgcaaa	tcatttcttc	acgaaaatgt	gctggcaaat	cagaagaaga	180
catgttgag	tgcttcattg	actccaagta	caaaaatgg	cgcccgacaa	ctgaggccga	240
ggctactgg	ctgcttattg	cggtctctct	tgcagggcag	cacaccagtt	ctatcacttc	300
cgtgtggact	ggggcctacc	tcctcaccaa	caagaagtac	ctctctgctg	tctctaata	360
acagaagcac	ctgatggaga	agcatgggaa	caatgttgat	catgatgttc	tttctgaaat	420
ggatgtcctg	tatcgggtcca	tc				442

<210> 129

<211> 392

<212> DNA

<213> Eucalyptus grandis

<400> 129

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gagaagcatg	ggaacaatgt	tgatcatgat	gttctttctg	aaatggatgt	cctgtatcgg	120
tccatcaagg	aagcactgag	acttcaccca	cctctaatta	tgctgctccg	aagctcgcac	180
agtgaattca	gtgtcaaaac	acgggatggc	aaggaatatg	agggtgggtga	agtctcagtg	240
cttccttgat	ggacccttga	ggcaaggaaa	gggtgcggca	aggcttttat	cactgcattc	300
aggtcgggtg	ccgtaatggg	cttccttctt	gctgcgaatg	gtcttctggg	cctttacatt	360
gccatcaacc	tattcaagat	ttacctatgg	gt			392

<210> 130

<211> 354

<212> DNA

<213> Eucalyptus grandis

<400> 130

gttgatttcg	acgtcgatta	caccatcagg	caagagcagt	ttcgggtttt	tactgaggct	60
ctgaggatta	ataagctcaa	ggggatgtgc	aatcagatgg	ttatggaagc	ggaggactac	120
ttctcaaaat	ggggagatag	tggcgagggtg	gacctaaagt	atgagcttga	gcacttgacc	180
atattgacag	cgagcagatg	tcttttgggt	cgagagggttc	gtgagaagct	ctttgatgat	240
gtgtcagccc	tattccacga	ccttgacaat	ggaatgctac	cgatcagtgt	catcttcccc	300
tacctgcccc	tcccagctca	ccatcgctcg	gataaggctc	ggaagaagct	tgct	354

<210> 131

<211> 442

<212> DNA

<213> Eucalyptus grandis

<400> 131

cttccgtgag	aagaagggtc	ttggagatga	caactggcag	atgtctggat	gggttgccct	60
------------	------------	------------	------------	------------	------------	----

tggatggatt	tgattatgga	tccatccttg	gccaatgctg	tgaactgcct	attgggtatg	120
tgagattcc	tgtgggtggt	gcagggcctc	ttctgcttga	tggaattgaa	aacatgggtc	180
ccatggccac	taccgagggc	tgcttgggtg	ccagcaccaa	cagaggttgt	aaggccattc	240
atatgtctgg	gggtgctaca	agcgttcttc	ttagagatgg	catgaccagg	gctcctgtcg	300
ttcgattccc	cactgctagg	agagctgcac	aactcaagtt	ttacttggaa	gccccataaa	360
ctacgaaagc	ttgtctctca	tcttcaacag	caccagcaa	ggtttgccag	gcttgcaaa	420
gaattcaagt	gcgccaatt	gg				442

<210> 132

<211> 984

<212> DNA

<213> Eucalyptus grandis

<400> 132

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agaagaagg	ctttggagat	tagaactggg	agatgtctgg	atgggttgcc	cttggatgga	120
tttgattatg	gatccattct	cggccaatgc	tgtgaactgc	ctgttgggtg	tgtgcagatc	180
cctgtgggtg	ttgtagggcc	tcttctgctt	gatggccttg	aaaacatggt	tcccatggcc	240
accaccgagg	gctgcctggt	ggccagcgct	aacagaggtt	gtaaggccat	tcatatgtct	300
gggtgtgcta	caagcgttct	tcttagagat	ggcatgacca	gagctcctgt	agttcgattc	360
cccactgccg	agagagctgc	acatctcaag	tcttacttgg	aacatcccaa	gaacttcgac	420
agcttgtctc	tcatcttcaa	cagcacaagc	agatttgcaa	ggctgcaaac	catcaagtgt	480
gcaattgcag	ggaggaatct	gtacataaga	ttttcctgct	tactggaga	tgccatggga	540
atgaatatgg	tgtccaagg	tgtgcagaat	gttttagact	tccttcagaa	tgaaaatcct	600
gatattggatg	ttattgctgt	ttctgtgta	ttctgtgccc	acaagaaacc	cacagctgtg	660
aactggattg	aagggcgtgg	aaaatccgta	gtttgcgagg	caattatcac	tgaagcagtt	720
gttagcaagg	ttttgaagac	caccgttcca	gctttgttag	aattgaacat	gctcaagaat	780
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gtctcagctg	tatttattgc	aacagggtcaa	gatcctgccc	agaacattga	gagctctcat	900
tgtattacga	tgatggaggc	atccaacgat	ggaaaggatc	ttcatgtatc	agtcaccatg	960
ccttgtattg	aggttggaaa	cagt				984

<210> 133

<211> 527

<212> DNA

<213> Eucalyptus grandis

<400> 133

ctcttggggg	agactgcagg	agagctgctt	cggtgagaag	aagggtcttg	gagatgacaa	60
ctggcagatg	tctggatggg	ttgcccttgg	atggatttga	ttatggatcc	atcctcgggc	120
aatgctgtga	attgcctgtt	ggatatgtgc	agattcctgt	gggtgttgca	gggcctcttc	180
tgctcgatgg	ctttgaaatc	atggttccga	tggccaccac	cgagggtgct	ctggtggcca	240
gcaccaacag	aggttgtaag	gccattcata	tgtctggggg	tgctacaagc	gttcttctta	300
gagatggcat	gaccagggct	cctgttgttc	gattctccac	tgctaggaga	gctgcacaac	360
tcaagtttta	cttggaaacat	ccaataaact	acaaaagctt	gtctctcatc	ttcaacagca	420
ccagcagatt	tgccaggctg	caagggaatca	agtgcgcaat	tgagggaagg	aatctgtaca	480
tgagggtttg	ttgttccact	ggagatgcca	tgggggatga	atatggg		527

<210> 134

<211> 965

<212> DNA

<213> Pinus radiata

<400> 134

aaacaaccag	ccgagttcga	aggtgggtcaa	cccacgtgta	ccatcatggc	acgtaagtag	60
atggcgaaaa	tggtggaagg	ccagttgtag	tgaccagggg	aatttaaaat	tattaatacg	120
ggactataaa	taactagcag	caatctctgg	cttttccact	gcattcatat	cgggagtttg	180
gggggattcc	aaaggatttc	ttcctctctc	tctccagtc	tcaatggatg	gccttgtgct	240
gaggaaaaa	agaaaattta	gtggagaaat	ggattgttgc	tccaacaaga	agatggagga	300
ttgtatggag	agttgtgggt	ctggattttc	tgggactgga	aagaagatga	aaaattcaag	360
gacattggca	tctgatgcct	tgccattgcc	tgtgggacta	accaacaagg	ttttctttat	420

cttggttttc	actgcttcct	atthttctgat	gaggagatgg	agggaaaaga	ttaggacttc	480
aacgccccct	catgtgctga	gcttagggga	gttggtcgcc	attgtggctc	agcttgcttc	540
attcatatat	ttgcttgga	tctttggcat	cgattatgtc	cagaatttca	tactggggg	600
caatgatgat	gatgatgcga	gggaagacga	taaactgagg	agccctgttc	ccaaggaagc	660
agttgcaatt	aggcccagtg	ctccgcaagt	ccagctgaac	gggatttcgt	tgggggataa	720
taaagatgat	gatattgcag	cagctgtctg	caatgggact	gtggcttctt	attctctcga	780
gtcgtctctt	ggggattgta	tgagatctgc	ccgggtgagg	aggaggtcct	tggagatgat	840
gactggcaga	tctttggatg	ggttgccctt	ggagggattc	gattatggat	ccattcttgg	900
ccaatgctgt	gaactgcctg	ttgggtatgt	gcagattcct	gtgggagttg	caggacctct	960
tcttc						965

<210> 135

<211> 503

<212> DNA

<213> Pinus radiata

<400> 135

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cagttggcat	cacaatctgc	ttgcttgaat	ttgatcggag	taaagggagc	gaatgtgcaa	120
tctcccgag	cgaatgctcg	gctcttggcc	aggattgtag	caggagcagt	tttggctgga	180
gagctctctc	tcatgtctgc	tttgggtgca	ggccagctgg	tcaagagcca	tatgaagtac	240
aacagatcaa	tcaaggatat	caaagcaatc	tcctgaacct	catggctcct	agaatccaag	300
aaagtcagca	tgggttttct	ccattgcgta	tctttactat	agcaatagac	ttatttgatc	360
aagctaggg	cctccaaaag	aaagtttcgt	acctgttacc	tgttttgcat	tgcataatgtt	420
atttgatcag	ctggggctct	ccaaaaggaa	gtttcctacc	tgctttcaat	tgcataatgtt	480
atttgatatt	catccaagtt	cta				503

<210> 136

<211> 563

<212> DNA

<213> Pinus radiata

<400> 136

ctcttttgaa	atacacacag	gcaagtcagc	agacatatcc	agagctcaat	cggcatatac	60
acagcaaac	aacaacatat	ttacaagcag	taaaatccac	cctgtagtga	tagtcccagg	120
cacaggagga	aatcagggtt	aagcaaggct	aactgcagac	tataaaccca	gtgggctggt	180
gtgcagaagg	tgaattggg	agagggagtg	gttcagaata	tggtttgatg	ttcctgtcgt	240
tcttctcca	ttgacacaa	gctttgctga	cagaataaag	ttggtctatg	atccccacac	300
agatgaatac	tacaatgccc	ctgggggtgga	gactagagtg	ccttattttg	gttcaacaga	360
aggaatgaag	taccttgatc	cctgcttcaa	gtatataacg	ccatacatgt	catccttggt	420
gaaatctctt	gaggatgttg	gatatgttga	cggtaaatcc	ctttttgggtg	caccctatga	480
tttcggttac	ggtcctggaa	caaagtcctc	ttctgtgggg	gcaaagtatt	tggaaaatct	540
gagaaaattg	gtggaggagg	cgt				563

<210> 137

<211> 354

<212> DNA

<213> Pinus radiata

<400> 137

ctcagcactt	atcataggca	gctttatctt	ttgcatcttt	ttatatatta	agggacatgt	60
tgcaccgtct	tcgactgatt	caggctcctc	tggaaatgta	gttattgatt	tctattgggg	120
tatggagctt	tatcctcgaa	taggtaaaaa	ctttgacatc	aaggctcttca	caaattgtcg	180
gtttggaatg	atgtcttggg	cagttcttgc	agtaacatac	agcataaaac	agtatgaaga	240
gtatggaaga	gtagcggatt	ccatgttagt	aagcagtata	ttgatgggtg	tgtatgtaac	300
aaaagtctct	cttgtgggaa	tctggctact	ggaacaccat	ggatataact	catg	354

<210> 138

<211> 631

<212> DNA

<213> Pinus radiata

<400> 138

ttcgcagttg	tgggacctct	gcagctgaca	tcgtatcccc	tgatcaagct	tgtgggtatc	60
agaacaggcc	tgccgttgcc	ttccctgtgg	gaaatttttg	cgcagcttgc	agtttatttc	120
atgggtgaag	actatggcaa	ctattggata	cacaggtggc	tacattgcaa	atggggctat	180
gagaagatcc	atcatgttca	ccatgagttc	actgctccaa	tgggttttgc	tgtcccatat	240
gcacattggt	cagaggtggt	gatattgggg	atccctacgt	ttgtcggacc	ggcaattgct	300
ccaggacaca	tgattacatt	ctgggtgctgg	gttgtgctgc	gccaagtggg	agcgattgaa	360
actcacagcg	gatatgactt	tccgtggact	cttaccaaa	taattccttt	ctatggaggc	420
gcggagtatc	atgactacca	tcattatgta	ggaggacaaa	gtcaaagcaa	ctttgcctca	480
gtgttcacat	actgtgatta	tttatacggg	actgataagg	gttaccgcta	tcgtaaggag	540
catcttttga	aggcacgtga	gtttgaatat	aggttaaagc	agatgatttt	aagaaagaaa	600
acggcaatgg	agcagtttca	gataagtttg	t			631

<210> 139

<211> 362

<212> DNA

<213> Pinus radiata

<400> 139

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ggcacattgt	ggttatgatt	ttccttggag	cattttcaaag	ctattcccgt	tgtatggagg	120
agctgatttc	catgattatc	atcatcgact	gctctatata	aagtctggga	attactcatc	180
gactttcact	tacatggact	ggttatttgg	gactgataaa	gggtaccgga	agctaaaaag	240
tctccagaaa	gattctaaat	gataacccaa	gagtgccatc	aaacatttgt	gtatgtgtgt	300
atcaattggt	tgaaggaaga	gaacagcaca	gacaggccta	gatcactctc	agtgaattcc	360
ag						362

<210> 140

<211> 504

<212> DNA

<213> Pinus radiata

<400> 140

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tcaccagatt	gggatttgtc	atggcgacat	tgggtgaaaag	aggctggctg	tatctgatca	120
caaatttcac	tgattttcaa	ctggcttcca	taggcagttt	tcttcttcat	gagagcatct	180
tctacttgct	tgggcttcct	ttcatattac	ttgagactac	aggcttggtg	agcaagtaca	240
aaattcagag	caagacgaac	acagttgctg	cacaagaaaa	atgtattact	cgactgctgc	300
tatatcattt	ttgtgtcaac	ctgccagtca	tgggtggttc	ctatcctgta	ttcagattta	360
tgggcattgac	aagcgtgcta	ccactaccat	cctggaaaag	agttgtatcc	caactggttt	420
gttatttcat	tttggaggat	tttgttttct	actggggcca	cagaatttta	cattcaaaat	480
ggctgtacaa	gcatgttcac	agtg				504

<210> 141

<211> 1293

<212> DNA

<213> Pinus radiata

<400> 141

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gtaaactatt	cctgagaagt	gtgtctgtta	cacaattctc	aaatatcatt	gatcttcagg	120
attttggatc	acatctgaga	acccaggat	gggagaagag	ttgcagacat	ggatattaat	180
ggtcactgct	agagctccta	caaatatagc	agtgatcaag	tactggggga	aaagagatga	240
aaagctgac	cttcccatca	atgacagcat	cagctttact	ttggatccag	accatctgtc	300
agccacaacc	actgtagcag	ttagcccatc	attcacatct	gatagaatgt	ggctcaacgg	360
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gggaaatgat	gtagtggacg	agaagaaggg	aattgttata	aggaaagagg	attggcagag	480
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ggctgctgga	ttcgcttgct	tagtttatgg	tctggcaaaa	ttaatggacg	tcaaggaaaa	600
atatcagggg	gaactttcag	ccattgccc	cagaggttca	gggagtgcac	gccgtagcct	660

ttatggtgga	gtggtaaaat	ggatgatggg	aaaggaaacc	gatggaagt	acagcattgc	720
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aattgcaaat	catgattttg	gatcctttgc	gaagattact	tgtgcagaca	gtaaccagtt	960
ccatgcagtt	tgcttgaca	catctcctcc	aatattctac	atgaatgaca	catcacacag	1020
gattattaac	tgtattgaaa	gatggaatcg	gtctgaagga	actccacagg	ttgcatatac	1080
ttttgatgct	gggtccaaatg	cagtaatgta	cgcacctaat	aggaaagtgtg	cagggtcacct	1140
tcttcagcga	ctgcttttct	atcttcctcc	ggactccagc	aaaacattgt	caagttatgt	1200
gataggcgac	acctcaatac	taggagaaat	cgcggtggac	tcaatgaagg	atgttgaatc	1260
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<210> 142

<211> 389

<212> DNA

<213> Pinus radiata

<400> 142

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actattcctg	agaagtgggt	ctgttacaca	attctcaaac	atcattgata	ttcaggattt	120
tggtcacat	ctgagaaccc	aggtatggga	gaagagtgtc	agacatggat	attaatgggtc	180
actgcaagag	ctctacaaa	tatagcagt	atcaagtact	gggggaaaag	agatgaaaag	240
ctgatccttc	ccatcaatga	cagcatcagc	tttactttgg	atccagacca	tctgtcagcc	300
acaaccactg	tagcagttag	cccatcattc	acatctgata	gaatgtgggt	caacggcaag	360
gaggtctctc	ttggagggga	gagatatca				389

<210> 143

<211> 693

<212> DNA

<213> Pinus radiata

<400> 143

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gaacagagaa	gggttaaca	cgatattgcc	tgaggaccaa	gtctataaga	tagaccagga	120
tagctatgcc	tctcactctt	ctattggcga	atacatgggc	atcgtctgcc	atagtttcca	180
ggaggggtatc	gctctttgtg	gcttgctcaa	caactgtagt	ctctcgttca	ttcagtaaga	240
gctgctccgg	tgctataccc	cgggaagccc	aatctgtctc	tcccgcactc	actgggagca	300
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tcaatacaaaa	atatgaattg	cttcttcagc	aacgttctgc	aacaaagggtg	acattccctt	600
tggtatggac	aaatacctgc	tgcagccatc	ctctctacag	ggagtctgag	ctcattgagg	660
agaacaattt	agggtcagaa	atgcagccca	aag			693

<210> 144

<211> 385

<212> DNA

<213> Pinus radiata

<400> 144

cgctgcagg	tcgacactag	tggtccaaa	gaagaactgg	tgtgatggca	ggaattccag	60
tcctaaggcc	atcttgcatc	tggttgcttt	cagtctacat	gctgcacatt	gtagctgcag	120
tagcttcacc	aaggctaggt	agaagcagct	tcccaggggg	tttcaaat	ggtgcagggt	180
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atacattctc	ccacactcca	ggtaaaatcg	ctgatgggaa	gaatggggat	gttgcatgag	300
atcaatacca	ccgttataag	gaagatgtgc	agcttctcaa	atacatggga	atggacgtct	360
atcgtttctc	tatctcctgg	tcacg				385

<210> 145

<211> 385

<212> DNA

<213> Pinus radiata

<400> 145

aaggccccag	catttgggat	acattctccc	acactccagg	taaaatcgct	gatggaaaga	60
atggggacgt	tgacgtagac	caataccacc	gttataagga	agatgtgcag	cttctcaaaa	120
acatgggaat	ggacgtctat	cgtttctcta	tctctgggtc	acgcataatt	cctaaggggt	180
cgccaagaca	cggaccagtc	aataaagtgg	gaatcgttta	ttacaataat	ttcatcaacg	240
agctgctcag	gaatgggtata	gagccttttg	tcacactgtt	tcactggggac	atgccacaag	300
ctctggaaga	tgagtacggg	ggattccgta	acaaaagagt	cgtggaggac	tttaacatat	360
ttgcagaagc	atgctttcga	gcctt				385

<210> 146

<211> 546

<212> DNA

<213> Eucalyptus grandis

<400> 146

ctccccctgtc	cttttctctc	tcccttcatt	aattctctct	tccgagatct	gatttttccct	60
cacttttccc	agaaaataat	ccccccgatc	tccccccggg	aattcccccc	cggccgttcg	120
attccggcgc	gcgctccggc	gatcgctcgc	tcgctcgcta	gccggttctt	ctctcgctcg	180
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tcgcggtgat	caagtactgg	gggaagcggg	acgagtcctt	catcttgccc	gtgaacgaca	300
gcacagcgt	gaccctggat	cccgggcacc	tctgcaccac	caccaccgtc	gccgtcagcc	360
ccgccttcga	gcaggaccgc	atgtgggtca	atggcaagga	gatatctctt	tctggagata	420
gatttcagag	ttgtttgaga	gaaattcgag	cccgtgctac	tgacgttgag	aataaggaaa	480
aaggaattaa	aatttcaaag	aaagattggg	agaaaactgca	cctccacatt	tctttcttta	540
catttc						546

<210> 147

<211> 786

<212> DNA

<213> Pinus radiata

<400> 147

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ggctctctcc	aggtacctgg	ttcgcttcc	ctgcatgttt	tttagacca	tagtttccc	120
acttacggaa	tttggttag	aattaggccc	tgcaaaagtt	ttatagcttc	ctctggggta	180
acggtagctt	acagggttga	attcgttgga	gcacgcgtga	gaagctacag	acatgagtag	240
caatggcaac	gggcaaaaac	aaggaggggg	ctttttcgcc	gccttcgcct	cgggcctctc	300
taatttcgga	agcgcgatgc	acaaatcggt	taacagcttc	atgggatatg	aggggttaga	360
agtagtcaat	cctgaaggcg	gtcaggatga	tgacagggag	gaagctcatc	gaggtagatg	420
gcggaaagag	gaccaggata	gttattggaa	aatgatgcaa	aaatatattg	gagcagatgt	480
cacctccatg	gtgacacttc	cagtcattat	ctttgagcct	atgacgatgc	ttcagaagag	540
tgctgagtta	atggagtaca	cttatttgct	tgacatggca	gatgagtgtg	aagatcccta	600
tctcaagatg	gcttatgcag	catcatgggc	aatttctgtc	tatcctgcat	accagaggag	660
ttggaagccc	tttaacccta	ttcttgga	aacttatgaa	atgggtcaatc	atggagggat	720
cacatttatc	gcagagcagg	tcagccacca	tcctccatgg	gctcagccta	tgccagaaat	780
gacatt						786

<210> 148

<211> 1748

<212> DNA

<213> Pinus radiata

<400> 148

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cgccttcgcc	tcgggectct	ctaatttcgg	aagcgcgatg	cacaaatcgg	ttaacagctt	180
catgggatat	gagggtttag	aagtagtcaa	tcctgaaggc	ggtcaggatg	atgcagagga	240
ggaagctcat	cgaggtagat	ggcgaaaga	ggaccaggat	agttattgga	aaatgatgca	300

aaaatatatt	ggagcagatg	tcacctccat	ggtgacactt	ccagtcatta	tctttgagcc	360
tatgacgatg	cttcagaaga	gtgctgagtt	aatggagtag	acttatttgc	ttgacatggc	420
agatgagtgt	gaagatccct	atctcaagat	ggcttatgca	gcacatggg	caatttctgt	480
ctatcctgca	taccagagga	gttggaaagcc	ctttaaccct	attcttggag	aaacttatga	540
aatggtcaat	catggaggga	tcacattttat	cgcagagcag	gtcagccacc	atcctccaat	600
gggctcagcc	tatgcagaaa	atgaacattt	tacatacagt	ctgtcctcaa	aagtaaaaaac	660
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ttatgtcatc	ttcattttgc	taatgtattt	tgaactcaag	ttaacagacg	atggaaacaa	1620
acttcctgtc	catgagttga	atgaaatttt	aaaagataat	ggaagctgcc	tccatgtgat	1680
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aaaaaaaa						1748

<210> 149

<211> 428

<212> DNA

<213> Pinus radiata

<400> 149

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tacagacatg	agtagcaatg	gcaacgggca	aaaacaagga	gggggctttt	tcgccgcctt	180
cgctcggggc	ctctctaatt	tcggaagcgc	gatgcacaaa	tcgggttaaca	gcttcatggg	240
atatgagggg	tagaagtagt	caatcctgaa	ggcggtcagg	atgatgcaga	ggaggaagct	300
catcgaggta	gatggcggaa	agaggaccag	gatagttatt	ggaaaatgat	gcaaaaatat	360
attggagcag	atgtcacctc	catggtgaca	cttcagtc	ttatctttga	gcctatgacg	420
atgcttca						428

<210> 150

<211> 419

<212> DNA

<213> Pinus radiata

<400> 150

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atttcatgtc	gcatectgat	ttgatcttcg	gggctgagaa	tagcaatgat	cctgaagaac	180
gattcatgcg	cgtactgtca	tactatttgg	ctgggttgga	tattaagcca	aaaggcgtca	240
aaaaaccgta	caatccagtt	cttggcgagt	ttttccgctg	tagatatgac	tattcgaata	300
atacacaagg	tttttatatt	gctgaacaag	tctctcatca	tccccccatt	tctgcatttt	360
tctacatttc	tcttgccaac	cgcgtgagca	tcatcgggga	actaagacca	aagtcaaaag	419

<210> 151

<211> 401

<212> DNA

<213> Eucalyptus grandis

<400> 151

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tcattgcttg	ccattgcgaa	ggaaggggtt	ggaaattctg	ggctgacagc	aatttaagga	180
caaaattttg	gggacaatct	attcagcttg	atcctgtggg	agcacttacc	cttgagtttg	240
atgatggcga	gattttttcaa	tggaataagg	taacaactag	catcaacaat	cttatcattg	300
gaaaagttta	ctgtgatcat	catgggtgtca	tgaatataca	tggaaccac	caatattcat	360
gcaaattgaa	gttcaaggag	ccatctattc	ttgccgaact	c		401

<210> 152

<211> 349

<212> DNA

<213> Eucalyptus grandis

<400> 152

cgcacatgcg	attgggtctat	gcgggcatca	tgggctatat	cagtctatta	tgccctatcaa	60
aggacgtgga	agcctttcaa	tcctattctt	ggggagactt	atgaactggc	aaatcatggc	120
ggtattactt	ttattgctga	gcagggtctgt	catcatcctc	caatgagtgc	cgggcatgcy	180
gaaaatgatc	attttaccgta	tgatgtgaca	tcaaaattaa	aaaccaaatt	cttaggggaa	240
tctgttgatg	tttatcctgt	aggaagaaca	cgtgtcactc	ttaaacgaga	tggtgtgggt	300
ttagatttgg	tgccaccccc	aacaaaggtt	aacaacctga	tttttgac		349

<210> 153

<211> 533

<212> DNA

<213> Eucalyptus grandis

<400> 153

ctctcgttta	cgtcctgata	gatatgcact	tgagccgggt	gaccttccta	aagctggtgc	60
tgaaaagagc	agcttggagg	aaaggcaaa	aggagagaaa	aagaaccgag	aaatgaaagg	120
ccagaaattc	actccaaggt	ggtttgatct	gactgacgaa	attagtccca	caccttgggg	180
cgatttgga	gtgtaccgct	acaatggaaa	gtatactgaa	catcgggctg	ttgtagacag	240
tctagacacc	atcgaagagt	ctgacattca	atcaactgag	ttcaatccct	ggcagtagca	300
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acctcctttt	tcttacgtag	tgcccaatgt	atatcatagt	tgatcgtctt	gaatgcgatt	420
agcatgttat	tcgccccttc	gttctttctac	tcagcatttt	tttattcata	ggagatcgta	480
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<210> 154

<211> 354

<212> DNA

<213> Pinus radiata

<400> 154

ggttcttcga	ggcctcgaca	ctggttgagga	tgatacaagc	attccattgg	atacaaagct	60
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agtcgagcac	tacaaagaac	tgatggaaaa	atttcatcat	gtttcaacta	ccttttttacg	180
gcttggaagg	ggatatcagg	aagcaattga	agaaataact	aagaagatgg	gtgctgggat	240
ggcaaaattt	atctgcaaag	aggttgaatc	agtggaggac	tatgatgaat	attgccatta	300
tgctgcagga	ctagttggat	ttggtttgtc	acgactcttt	catgcagctc	agct	354

<210> 155

<211> 675

<212> DNA

<213> Pinus radiata

<400> 155

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tcaaccagca	catcgactga	tatcgtggtc	tacaatggag	aatcatacgg	tggtgattgc	180
ggcagccatt	agctttgttt	ctgtattatt	gtcgtattat	atagttttga	gcagggtggaa	240
gcgcagatcc	aacggattac	ggggaatata	gagcaaaagt	ttcgaaaagt	caacagatga	300

caatggcatt	gccatcgaag	ctgctggagg	aacggatggt	atcatcgtgg	gagcaggagt	360
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gaaattgatt	gagctggggac	ttcaagattg	tgttgaaggga	attgatgccc	agagtatatt	540
tggggatgct	ttattcaagg	aaggaaaaga	tactaaagtg	gcatatccgt	tagaaaacca	600
ccatgcagat	agagctggaa	ggagtttcca	caatggacgc	ttcatccagc	gcatgcgggga	660
aaaggctgct	tcact					675

<210> 156

<211> 373

<212> DNA

<213> Pinus radiata

<400> 156

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atctatacgc	ctcaaccagc	acatcgactg	atatcgtggt	ctacaatgga	gaatcatact	120
gtggcgattg	cggtagccat	tggctttgtt	tctgtattat	tgtcgtatta	tatagttttg	180
aacagggtgga	agcgcagatc	caacggatta	cggggaatac	agagcaaaag	tttcgagaag	240
tcaacagatg	acaatggcat	tgccatcgaa	gctgctggag	gaacggatgt	tatcatcgtg	300
ggagcaggag	tcgcggttc	ggctctggct	tacacacttg	gcaaggatgg	aagacgtata	360
catgtaattg	aga					373

<210> 157

<211> 522

<212> DNA

<213> Eucalyptus grandis

<400> 157

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gcaagagccg	agctccatcg	gcggaagctt	ctcgatctct	ctgctgtcgc	gtggcgagg	120
ttggtggagc	tgttaagccg	gatcggctct	tgatggacgg	tcagtacttg	gtcagtgggc	180
tcttggtctt	gttcctgggg	atcttcctgc	tgtacaaggg	gctcgggaag	cagaagagga	240
ggctgtccaa	gaagggtcgc	ggcgatgact	atgtgaagag	ctctgtggat	ggagggttgc	300
tgcccggcgg	cgctgatggg	agcaccgaca	tcgtcattgt	cggagcaggc	gtcgcgggtg	360
cggctctcgc	ttacgccctc	gggaaggatg	gacgtcgcgt	gcgtgtaatt	gagagggacc	420
tgacggagca	agatagaatt	gtcggcgagc	ttcttcaacc	aggaggttac	ctgaaattga	480
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<210> 158

<211> 898

<212> DNA

<213> Eucalyptus grandis

<400> 158

ctcgggtcga	agtataaacc	tcaggaagaa	tttgttgaat	ggattcaaaa	gggaacaaaa	60
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ataattaagg	cattaacgga	taccggacaa	agagggatag	ttggtcgagg	ttggggtgat	180
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ggcgataggg	tccaccaaag	aggccttggc	cctgcaccaa	taccaatctc	ccagctcagc	420
gtcgagaacc	tttcagatgc	cataagattt	atgcttcaac	ctgaggtaaa	gtctcaggca	480
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ttcatttcat	tcttatcagg	gtttggctga	ccattgtatt	cagcatagca	taagatttaa	840
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<210> 159

<211> 342

<212> DNA

<213> *Eucalyptus grandis*

<400> 159

ctcgataatt gccctcatga ctggcttttc ctgcgctgca gtgctgtggt acatcatgga	60
ggagctggta caaccgctgc cggctcttaaa gctgcgtgtc caacaacagt tgtacctttc	120
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<210> 160

<211> 582

<212> DNA

<213> *Pinus radiata*

<400> 160

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cttatgatcc cctaaccctt aaatcgtagc agtgaagcca ttaacgattt ttgcgggttc	120
agaaagattc actgaatcgc ttactaaaac tctgtttcag gaatggcaac aggaggagga	180
gcgttggatc tggcctcagg aatgggaggg aacattgaga aagaacaaat gctgaccgct	240
gttgaagagt acgaaaaata tcacatgtac tatggtgggtg atgaaggctc gagaaaatct	300
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gatgttggat gtgggattgg aggtccactg agagaaattg ctaggttcag tcggacttcg	540
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<210> 161

<211> 552

<212> DNA

<213> *Eucalyptus grandis*

<400> 161

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cctgcattgt cctcattctg ggcgggggtg ccacaatgtc gaaagcagga gcgatggatc	120
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taaacaacaa tgagtaccag ataacaaggg gaaaggaact aaaccgcatt gcaggcgtgg	540
acaagacatg cg	552

<210> 162

<211> 401

<212> DNA

<213> *Eucalyptus grandis*

<400> 162

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<210> 163

<211> 446

<212> DNA

<213> Eucalyptus grandis

<400> 163

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 aatatatctt gtaatggcgg agtttgggat atattggatg cacagagagc tgcacgacat 180
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Y2K

<210> 164

<211> 823

<212> DNA

<213> Eucalyptus grandis

BUSINESS CONTINUITY PLAN

<400> 164

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DOCUMENT AND INFORMATION SERVICE
CENTRE

<211> 90

<212> PRT

<213> Eucalyptus grandis

17 TOOP ST

<400> 165

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 Arg Asp Lys Ala Arg Lys Lys Leu Ser Glu Ile Phe Ala Asn Ile Ile
 35 40 45
 Ser Ser Arg Lys Cys Ala Gly Lys Ser Glu Glu Asp Met Leu Gln Cys
 50 55 60
 Phe Ile Asp Ser Lys Tyr Lys Asn Gly Arg Pro Thr Thr Glu Ala Glu
 65 70 75 80
 Val Thr Gly Leu Leu Ile Ala Ala Leu Phe
 85 90

<210> 166

<211> 40

<212> PRT

<213> Eucalyptus grandis

PREPARED DECEMBER 1999 REVISION 7

<400> 166

Tyr Leu Leu Thr Asn Lys Lys Tyr Leu Ser Ala Val Ser Asn Glu Gln

1	5	10	15
Lys His Leu Met Glu Lys His Gly Asn Val Asp His Asp Val Leu			
20	25	30	
Ser Glu Met Asp Val Leu Tyr			
35			

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<212>	PRT
<213>	Eucalyptus grandis
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Arg Leu Val Phe Ala Glu Gly Glu Asp Gly Pro Tyr Leu Tyr Ser Thr	
1	5
Asn Asn Val Phe Ile Trp Glu Phe Asp Pro Glu Ala Gly	
20	25
Thr Ala Glu Glu Arg Ala Glu Val Glu Ala Ala Arg Gln His Phe Tyr	
35	40
Asp His Arg His Gln Val Lys Pro Cys Gly Asp Leu Leu Trp Arg Met	
50	55
Gln Phe Leu Arg Glu Lys Glu Phe Lys Gln Thr Ile Pro Pro Val Arg	
65	70
Val Glu Asp Gly Glu Glu Ile Thr Tyr Asp Lys Ala Ser Thr Ala Leu	
	90
Lys Arg Ala Val His Phe Phe Ser Ala Leu Gln Ala Ser Asp Gly His	
	105
Trp Pro Ala Glu Asn Ala Gly Pro Leu Phe Phe Leu Pro Pro Leu Val	
	120
Met Cys Val Tyr Ile Thr Gly His Leu Asp Ala Val Phe Pro Ala Glu	
	135
His Arg Lys Glu Ile Leu Arg Tyr Ile Tyr Asn His Gln Asn Glu Asp	
145	150
Gly Gly Trp Gly Leu His Ile	
	155
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Met Asp Asp Ile Val Ser His Glu Phe Glu Gln Lys Arg Gly His Val	
1	10
Val Ser Ala Val Glu Leu Leu Ile Lys Tyr Arg Gly Val Ser Glu Gln	
	25
Glu Ala Val Glu Glu Leu Gln Lys Arg Val Ile Asp Ala Trp Lys Asp	
	40
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Thr Asn Glu Glu Phe Leu Arg Pro Ile Ala Val Pro Met Pro Ile Leu	
50	55
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Thr Arg Val Leu Asn Leu Ser Arg Val Ile Asp Val Leu Tyr Ser Asp	
65	70
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Gly Asp Asn Tyr Thr His Ser Glu Thr	
	85
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11.	REPORT ON TESTING
Met Glu Asp Asp Arg Asp Arg Gly Leu Leu Tyr Asp Ser Asp Pro Pro	

1	5	10	15
Ser	Pro	Ser	Leu
20	25	30	
Thr	Phe	Phe	Asp
35	40	45	
His	Met	Leu	Pro
50	55	60	
Leu	Ala	Tyr	Pro
65	70	75	
Arg	Glu	Arg	His
80	85	90	
His	Met	Leu	Pro
95	100	105	
His	Met	Leu	Pro
110	115	120	
His	Met	Leu	Pro
125	130	135	
His	Met	Leu	Pro
140	145	150	
His	Met	Leu	Pro
155	160	165	
His	Met	Leu	Pro
170	175	180	
His	Met	Leu	Pro
185	190	195	
His	Met	Leu	Pro
200	205	210	
His	Met	Leu	Pro
215	220	225	
His	Met	Leu	Pro
230	235	240	
His	Met	Leu	Pro
245	250	255	
His	Met	Leu	Pro
260	265	270	
His	Met	Leu	Pro
275	280	285	
His	Met	Leu	Pro
290	295	300	
His	Met	Leu	Pro
305	310	315	
His	Met	Leu	Pro
320	325	330	
His	Met	Leu	Pro
335	340	345	
His	Met	Leu	Pro
350	355	360	
His	Met	Leu	Pro
365	370	375	
His	Met	Leu	Pro
380	385	390	
His	Met	Leu	Pro
395	400	405	
His	Met	Leu	Pro
410	415	420	
His	Met	Leu	Pro
425	430	435	
His	Met	Leu	Pro
440	445	450	
His	Met	Leu	Pro
455	460	465	
His	Met	Leu	Pro
470	475	480	
His	Met	Leu	Pro
485	490	495	
His	Met	Leu	Pro
500	505	510	
His	Met	Leu	Pro
515	520	525	
His	Met	Leu	Pro
530	535	540	
His	Met	Leu	Pro
545	550	555	
His	Met	Leu	Pro
560	565	570	
His	Met	Leu	Pro
575	580	585	
His	Met	Leu	Pro
590	595	600	
His	Met	Leu	Pro
605	610	615	
His	Met	Leu	Pro
620	625	630	
His	Met	Leu	Pro
635	640	645	
His	Met	Leu	Pro
650	655	660	
His	Met	Leu	Pro
665	670	675	
His	Met	Leu	Pro
680	685	690	
His	Met	Leu	Pro
695	700	705	
His	Met	Leu	Pro
710	715	720	
His	Met	Leu	Pro
725	730	735	
His	Met	Leu	Pro
740	745	750	
His	Met	Leu	Pro
755	760	765	
His	Met	Leu	Pro
770	775	780	
His	Met	Leu	Pro
785	790	795	
His	Met	Leu	Pro
800	805	810	
His	Met	Leu	Pro
815	820	825	
His	Met	Leu	Pro
830	835	840	
His	Met	Leu	Pro
845	850	855	
His	Met	Leu	Pro
860	865	870	
His	Met	Leu	Pro
875	880	885	
His	Met	Leu	Pro
890	895	900	
His	Met	Leu	Pro
905	910	915	
His	Met	Leu	Pro
920	925	930	
His	Met	Leu	Pro
935	940	945	
His	Met	Leu	Pro
950	955	960	
His	Met	Leu	Pro
965	970	975	
His	Met	Leu	Pro
980	985	990	
His	Met	Leu	Pro
995	1000	1005	
His	Met	Leu	Pro

Asp Ser Glu Pro

130

	Service Centre	Corporate Services
Person responsible for overall management of response	Janet Dobbie	Gary Jones or Shirley Herewini
Other Management Team Members (and roles)	Shirley Herewini Team Leader Records Gary Jones Team Leader Post Acceptance	
<400> 172 Arg Thr Leu Arg Leu His Gly Tyr Pro Val Ser Ser Asp Val Leu Lys	Sue Whiteman, Support Services Theresa King, Revenue & Lodgement	
1 Val Thr Leu Arg Leu His Gly Tyr Pro Val Ser Ser Asp Val Leu Lys	Leader: Janet Dobbie Members: 30	Ile Gln
Thr Gly Gly Glu Ile Arg Gly Val Leu Ser Thr Arg Ala Ser Leu	Shirley Herewini Gary Jones 45	
Val Ala Phe Pro Gly Glu Asn Val Leu Ser Thr Arg Ala Glu Ile Phe Ser	Sue Whiteman	
Recovery Team: 55 Thr Thr Tyr Leu Lys Glu Ala Ser Thr Val Pro Ile Ser Ser Ala	Leader: Janet Dobbie 60 Members: 75	
65 Ser Leu Ser Arg Glu Ile Glu Tyr Val Leu Glu Tyr Arg Trp Leu Thr	Shirley Herewini Gary Jones 95	
Asn Phe Pro Arg Leu Glu Ala Arg Asn Tyr Ser Asp Leu Phe Gly Asn	Sue Whiteman	
Other Key Staff: (see 3 of this group)	Tania McConnochie 110 Theresa King 125 Jenny Spaans 125 Margaret Newton 140 Joanne Sexton 140 John Apple 150 William Rodrigues	
Media Contacts	Janet Dobbie	Diane Imus

Key external Contacts:

Power	TransAlta	Phone: (04) 568 8800
Gas (not supplied to Toop St)	TransAlta	Phone: (04) 568 8800
Water/Effluent Floodings	Hutt City Council After Hours Emergency Service	Phone: (04) 567 2003
Suppliers	DX Post Haste Couriers	Phone: (04) 473 9510 Phone: (04) 499 2121
Others:	ABS (Cleaners)	Phone: (04) 564 3249
Val Ile Glu Tyr Gly Trp Leu Arg Thr Ser Thr Arg Leu Glu Ala Arg	Armourguard Security	Phone: (04) 478 1226
Asn Tyr Ile Asp Val Phe Gly Lys Asp Thr Ile Pro Cys Val Lys Thr	Monitoring Centre	Phone: (04) 478 1226
65 Thr Thr	Nedax Security	Phone: (04) 478 2836
<210> 174	Building Manager Melanie Lambert	Phone: (04) 499 9133 After Hours (04) 934 2552 or (025) 262 3146
<211> 141		
<212> PRT		

Emergency Services:

Fire	111 or Petone Fire Brigade (04) 568 6857
Ambulance	11 or Wellington Free Ambulance (04) 472 2999
Police	111 or Petone Police Station (04) 568 7335
Boisons Centre	Urgent (03) 474 7000 Non Urgent (03) 479 1200
Occupational Health Centre	OK Health Phone: (04) 568 7128
Accident and Medical Centre	Hutt Hospital Phone: (04) 566 6999
Lower Hutt Civil Defence	Brian Toomey or Bill Flemming Phone: (04) 570 6666

His Ser Leu Gln Gln Lys Glu Leu Lys Gln Leu Ser Arg Trp Trp Lys
 50 55 60
 Asp Ser Gly Phe Ser Gln Leu Thr Phe Thr Arg His Arg His Val Glu
 65 70 75 80
 Phe Tyr Thr Leu Ala Ser Cys Ile Ala Thr Glu Pro Lys His Ser Ala
 85 90 95

Phe Arg Leu Gly Phe Ala Lys Thr Cys Tyr Leu Gly Ile Val Leu Asp
 100 105 110
 Asp Ile Tyr Asp Thr Phe Gly Thr Met Glu Glu Leu Glu Phe Thr
 115 120 125

Ala Ser Thr Trp Asp Ser Ala Arg Glu Phe
 130 135 140

Scenario Presented Invoke Plan Decision

No access to Seaview area	No
No building access	Yes
Building access, no utilities	Yes
Building access, no IT	Yes
Building access, no IPOL	Yes
No Mail Delivery	Yes

<400> 175
 Leu Thr Asn Phe Pro Arg Leu Glu Ala Arg Asn Tyr Ile Asp Val Phe
 1 5 10 15
 Gly Ile Asn Thr Ser Val Arg Thr Met Glu Glu Leu Glu Phe Thr Ala
 20 25 30
 Thr Glu Cys Thr Phe Gly Thr Met Glu Glu Leu Glu Phe Thr Ala
 35 40 45
 Ser Leu Gln Gln Lys Glu Leu Lys Gln Leu Ser Arg Trp Trp Lys Asp
 50 55 60
 Ser Gly Phe Ser Arg Leu Thr Phe Thr Arg His Arg His Val Glu Phe
 65 70 75 80
 Tyr Thr Leu Ala Ser Cys Ile Ala Thr Glu Pro Lys His Ser Ala Phe
 85 90 95
 Arg Ile Tyr Thr Trp Asp Ser Ala Arg Glu Cys Leu Pro Glu Tyr
 100 105 110
 Ile Tyr Asn Thr Phe Gly Thr Met Glu Glu Leu Glu Phe Thr Ala
 115 120 125
 Ala Ile Lys Arg Trp Asp Pro Ser Ala Arg Glu Cys Leu Pro Glu Tyr
 130 135 140
 Met Lys Gly Ile Tyr Met Val Phe Tyr Asp Ala Leu Ile Lys Trp Leu
 145 150 155 160
 Glu Arg

3.1 No Access to Seaview
 In the event that Seaview suffers major infrastructure problems the Impact Assessment Team will:
 • Contact staff by phone tree
 • Leave recorded phone message on (04) 568 0744 for staff information and (04) 568 0720 for Clients information.

3.2 No Building Access
 In the event the Documents & Information Service Centre remains closed the Impact Assessment Team will:
 • Contact staff by phone tree
 • Leave recorded phone message on (04) 568 0744 for staff information and (04) 568 0720 for Clients information.

3.3 Building Access no Utilities
 If the building is open but there is no power, water or sewerage the Impact Assessment Team will:
 • If staff not present, contact by phone tree.
 • If staff already present, consider sending staff home.
 • If no power, the leader will contact Manager IT Operations.
 • Leave recorded phone message on (04) 568 0744 for staff information and (04) 568 0720 for Clients information.

3.4 Building Access no IT
 If all services are available except IT systems the Impact Assessment Team will:
 • Re-assign staff to other work.

3.5 Building Access but no IPOL
 If the IPOL system is not available the Impact Assessment Team will:
 • Re-assign staff to other work.

<212> PRT

<213> Eucalyptus grandis

<400> 177

Leu 1.6 Gln Phe Val Ala Asp Leu Lys Gly Glu Phe Leu Asn Arg

1 5 10 15

Lys If mail is unable to be delivered NZ Post will store the mail until collection can be organised
either by DX Courier or ourselves.Ser Ser Gln Leu Phe Cys Met Glu Asn Asp Gly Phe Thr His Ser His
35 40 45

Glu Thr Glu Ile Lys Glu His Val Lys Lys Ile Leu Phe Glu Pro Val

50 55 4. PLAN SUMMARY

Ala

65

4.1 Objectives

<210> 178

This plan outlines the steps necessary to determine the ability of the Document and Information
Service Centre to provide services to IPONZ and its external clients beyond 1 January 2000.

<213> Eucalyptus grandis

4.2 Before 24 December 1999

<400> 178

Leu Asp Cys Glu Pro Val Val Gln Lys Pro Lys Leu Val Asp Pro Val

Task	Responsibility	Target Date	Completed
1. Seek assurances from suppliers for Y2K compliance	Sue Whitman	13 July 1999	Yes
2. Test all software and hardware for Y2K compliance	IBB & Co	13 July 1999	Yes
3. Discuss with support services arrangements for alternative accommodation	Shirley Herewini	30 September 1999	Yes
4. Contact key suppliers to obtain and confirm contact names	Shirley Herewini Sue Whitman	31 October 1999	Yes
5. Distribute copies of BCP plan to all staff for home and work	Sue Whitman	15 November 1999	Yes
6. Inform clients and staff of holiday period telephone number on which messages will be left	Glenn Dobson	1 December 1999	Yes
7. Arrange necessary stationery for date stamping	Shirley Herewini	15 December 1999	Yes
8. Arrange storage for mail and cheques	Shirley Herewini	15 December 1999	Yes
9. Arrange van to be full of petrol.	Shirley Herewini	24 December 1999	

Lys Phe Phe Val Glu Asn Pro Ala Asn Phe Glu Ser Leu Ala Val Ile
180 185 190Phe 4.3 After 1 January 2000
195 200 205

Task	Responsibility	Target Date
1. Check that location is accessible, building secure, power on	Armourguard	12:30 am - Armourguard Security to contact Neville Harris (Diane Imus as backup to Neville)
2. If no power or access to office	Neville Harris to contact Mike Brosnahan	By 1am 1 Jan
3. BRB testing of IT LAN/WAN systems	Michael Brosnahan	5 Jan 16am (initial IT ck)
4. Joe Ryan (can you do a check)	Joe Ryan	5 Jan 11am 1 Jan
5. Full BRB test start by 1pm 1 Jan. MB to report to Kathryn McInteer by 3pm		

Asn Ala His Ala Ser Asn Ile Val Ala Ala Ile Phe Ile Ala Thr Gly	325	330	335
Gln Asn Val Glu	340	345	350
Glu Ala Ile Asn Asp Gly Lys Asp	355	360	365
Ser Val Glu Val Gly Thr Val Gly	370	375	380
Ser Ala Cys Leu Asn Leu Leu Gly	385	390	395
Ala Gly Ala Asn Ser Arg Leu Leu	405	410	415
Leu Ala Ala Glu Leu Ser Leu Met	420	425	430
Val Tyr Asn	435	440	445
Val Ser	450	455	460
Val Ser	465	470	475
Val Ser	480	485	490
Val Ser	495	500	505
Val Ser	510	515	520
Val Ser	525	530	535
Val Ser	540	545	550
Val Ser	555	560	565
Val Ser	570	575	580
Val Ser	585	590	595
Val Ser	600	605	610
Val Ser	615	620	625
Val Ser	630	635	640
Val Ser	645	650	655
Val Ser	660	665	670
Val Ser	675	680	685
Val Ser	690	695	700
Val Ser	705	710	715
Val Ser	720	725	730
Val Ser	735	740	745
Val Ser	750	755	760
Val Ser	765	770	775
Val Ser	780	785	790
Val Ser	795	800	805
Val Ser	810	815	820
Val Ser	825	830	835
Val Ser	840	845	850
Val Ser	855	860	865
Val Ser	870	875	880
Val Ser	885	890	895
Val Ser	900	905	910
Val Ser	915	920	925
Val Ser	930	935	940
Val Ser	945	950	955
Val Ser	960	965	970
Val Ser	975	980	985
Val Ser	990	995	1000

<210> 181
 <211> 81
 <212> PRT
 <213> Eucalyptus grandis

 <400> 181

Ser Ile Lys Cys Ala Ile Ala Gly Lys Asn Leu Tyr Leu Arg Phe Ser
 1 5 10 15
 Cys Ser Thr Gly Asp Ala Met Gly Met Asn Met Ile Ser Lys Gly Val
 20 25 30
 Gln Asn Val Met Asp Phe Leu Gln Lys Asp Phe Pro Asp Met Asp Val
 35 40 45 50 55 60

STAFFING AND SUPPORT

Met Gly Ile Ser Gly Asn Phe Cys Ser Asp Lys Lys Pro Ala Ala Val
 50 55 60

Asn Trp Ile Glu Gly Arg Gly Lys Ser Val Val Cys Glu Ala Val Ile
 65
 Lys The Impact Assessment Team is required to be on call to assess the situation after Computer Services BRB IT have completed testing on 2 January 2000. Leader, Janet Dobbie will be responsible for informing the rest of the Team.

Those on call on 1 January 2000 need only be contacted for a progress report on the outcome of IT services availability. They will be required to make preparations to return to Wellington if the need arises. PRT

<213> Pinus radiata

Person	Availability	Date & Time	Back-up Resource
Janet Dobbie	On call	Glu Glu Val Val Lys	Diane Imus
Sue Whiteman	On call	10 15	
Gary Jones	On call	Glu Leu Asn Met Leu	
Shirley Heregoni	On call	25 30	
Rest of DISC staff	On site unless otherwise advised	15 January 8am	Phe Asn
35	40	45	

Ala His Ala Ser Asn Ile Val Ser Ala Ile Tyr Ile Ala Thr Gly Gln
 50 55 60

Asp Pro Ala Gln Asn Val Glu Ser Ile Thr Met Met Glu
 65 70 75 80

Ala Val Asn Glu Gly Arg Asp Leu His Ile Ser Val Thr Met Pro Ser
 85 90 95

Ile Glu Val Thr Gly Ile Val Gly Gly Thr Gln Leu Ala Ser Gln Ser
 100 105 110

Ala The Impact Assessment Team are responsible for determining the operational status of DISC and the impact on IPONZ operations
 125

Gly Ala Asn Ala Arg Leu Leu Ala Thr Ile Val

On or by 2 January 2000 Janet Dobbie will check:

- Power is OK (Armourguard to report back to Neville).
- <210> 183 Access to IPOL is available - Mike Brosnahan by 4pm 1 Jan
- <211> 269 If IPOL or IT is not available BRB IT will be called to assess the problem and make an assessment call on how to proceed.
- If all IT and IPOL OK - Diane Imus to contact Janet and keep informed.

Met Asp Val Arg Arg Arg Gln Pro Lys Pro Pro Arg Pro Ala Ala Gly
 1 5 10 15
 Impact Assessment Team and staff using the telephone call tree.

Asp Pro Arg Arg Arg Gln Lys Ser Leu Arg Leu Pro Ala Pro Gly Val
 20 25 30 35 40 45 50 55 60
 As a result the Impact Assessment Team will determine the effects of any problems on

Asp Arg Arg Arg Arg Gln Pro Ser Pro Ser Pro Lys Ala Ser Asp Ala Leu
 60 65 70 75 80 85 90 95

Pro Leu Pro Leu Tyr Leu Thr Asn Ala Val Phe Phe Thr Leu Phe Phe
 50 55 60
 Ability to function, and then

Ser Val Ala Tyr Tyr Leu Leu His Arg Trp Arg Asp Lys Ile Arg Ser
 65 70 75 80 85 90 95
 Recovery Team

Ser Val Pro Leader Notify staff and clients the recorded message on phones. Ile Val
 90 95

Ser Leu Ile Ala Ser Phe Ile Tyr Leu Leu Gly Phe Phe Gly Ile Gly
 100 105 110

Phe Val Gln Phe Phe Ile Ala Ser Arg Ala Ser His Asp Ala Trp Asp Val
 115 120 125
 Implement a manual system of date stamping

Leu Asp Asp Glu Val Ala Val Gly Gly Asp Gly Phe Leu Pro Glu Asp
 If a manual system is required the Team will:

130 135 140
 Asp Gly Pro Pro Cys Ala Ala Ile Ala Cys Ala Pro Pro Lys Leu Ala
 145 1. Decide site to be used. 155 160
 Glu Arg Gln Note: The above information will already be on site as part of the pre 14 December 1999
 preparations. 165 170 175
 Val Val Lys Ser Val Thr Asp Gly Lys Ile Pro Ser Tyr Ser Leu Glu
 180 2. Assign Records personnel to the site to: 185 190
 Ser Met Leu Gly Asp Cys Lys Arg Ala Thr Ser Ile Arg Arg Glu Ala
 195 200 205
 Leu Gln Arg Met Thr Gly Arg Ser Ser Lys Gly Leu Pro Leu Glu Gly
 210 • Put all mail and cheques in date order and deposit in safe.
 Phe Asp Tyr Glu Ser Ile Leu Gly Gln Cys Cys Glu Met Pro Val Gly
 225 3. If in usual building and can work, cashiers will: 240
 Tyr Val Gln Ile Arg Trp Gly Ser Pro Val Arg Cys Cys Ser Thr Gly
 245 250
 Trp Ser Ile Pro Cys Trp Arg Trp Pro Arg Val Ala 255
 260
 • Write up abstracts for private applicants,
 • Balance and check all abstracts and cheques,
 <210> 264 All cheques and abstracts in date order and put in safe.
 <211> 279
 <212> PRT
 <213> Eucalyptus grandis

7. RETURN TO NORMAL

<400> 184
 Met Asp Val Ser Arg Pro Ser Lys Pro Ala Ala Ala Gly Ser
 1 5 10 15
 Ser The Recovery Team will declare a return to normal situation when:
 20 25 30
 Ala 1. Primary site, the building and Toop Street is usable. Phe Leu Tyr Leu Val
 2. IT link to Toop Street is working and, 45
 Asn 3. IPO is fully functional. All ops are the Thr Leu Leu Tyr Tyr Leu Leu
 50 55 60
 Ser Arg Trp Arg Glu Lys Ile Arg Ser Ala Ser Pro Leu His Val Leu
 65 70 75 80
 Ser Ala Pro Glu Leu Ala Ala Ble TEST AND PREVIEW Ala Ser Ser Val
 85 90 95
 Tyr Leu Leu Gly Phe Phe Gly Val Glu Phe Phe Gln Ser Leu Leu Leu
 100 105 110
 Arg The BCP Review Team will review the plan on a monthly basis to ensure all information
 continues to be correct. 120 125
 Val Ser Val Val Ala Glu Glu Ala Leu Lys Ala Pro Cys Gly Gln
 130 The plan will be discussed with all staff to ensure: 140
 Ala • All staff know whom to contact if necessary, 150
 145 • All staff are aware of the tasks expected of them before and after the event. 155 160
 Ala • That all pre-arrangements have been completed by the dates specified. 165 170
 175
 Ile • That all contact information is up to date. Val Ile Ala Ser Val Val Ala
 180 185 190
 Gly In the event this plan has been invoked the Impact Assessment Team will be required to conduct
 a Post Disaster Appraisal within 14 working days of return to normal. Comments and criticism
 Arg will be sought from all parties where appropriate to determine what improvements can be
 210 215 220
 Ser Leu Ala Gly Leu Pro Leu Glu Gly Leu Asp Tyr Ser Ile Leu
 225 230 235 240
 Gly Gln Cys Cys Glu Met Pro Val Gly Tyr Val Gln Ile Pro Val Gly
 245 250 255
 Ile Ala Gly Pro Leu Val Leu Asp Gly Arg Glu Tyr Ser Val Pro Met
 260 265 270
 Ala Thr Thr Glu Gly Cys Leu
 275

<210> 185

<211> 194

<212> PRT

<213> Eucalyptus grandis ² COMMUNICATION PLAN

Sue Whiteman will be responsible for ensuring copies of the plan are distributed via e-mail to all staff.

Met Val Arg Arg Arg Pro Pro Lys Pro Pro Leu Pro Ser Ala Ala
1 5 10 15

Arg Gly Gly Gly Arg Gly Pro Ala Ser Ser Ser Pro Pro Leu Glu Pro
20 25 30

Pro Lys Ala Ser Asp Ala Leu Pro Leu Pro Leu Tyr Leu Thr Asn Ala
A further copy will be distributed to all staff on the 22nd December 1999.

Val Phe Phe Thr Leu Phe Phe Ser Val Ala Tyr Tyr Leu Leu His Arg
Communicating to staff of developments - phone message on a particular number (04 568 0744)

Trp that they may require. Phone message will be updated by Janet Dobbie. Thr Leu
65 70 75 80

Pro Communicating to clients and BRB, MOU phone message (04 568 0720) to be changed by
Janet Dobbie. 85 90 95

Leu Gly Phe Phe Gly Ile Asp Phe Val Gln Thr Phe Ile Ala Arg Ala
Poster on door if the building is not in use. 100 105 110

Ser His Asp Ala Trp Glu Asp Leu Asp Asp Asp Val Asn Arg Gly Phe
Media release (if necessary) to be done by the Ministry Communications Unit ONLY
115 120 125

Gly Cys Thr Asp Ile Val Ala Pro Leu Pro Lys Ser Gly Asp Pro Ala
130 135 140

Pro Val Ile Ser Ala Leu Ser Ser Ala Glu Asp Glu Glu Ile Val Lys
145 150 155 160

Ser Val Val Asp Gly Thr Ile Pro Ser Tyr Ser Leu Glu Ser Lys Leu
165 170 175

10. APPENDIX

Gly Post Contact Details Ala Ala Phe Val Arg Arg Glu Ala Leu Gln Arg

Name	Position	Contact Number
Janet Dobbie 138 Cockayne Road Ngairi Wellington	Manager, Document & Information Service Centre	(04) 568 0720 work (04) 479 7539 home 021 362 898 mobile
Gary Jones 28 Exploration Way Whitby	Team Leader Post Acceptance	(04) 568 0726 work (04) 234 1400 home
Shirley Pieroni PO Box 110 Shelly Bay	Team Leader Records Ser Leu Pro Ser Asp Phe	(04) 568 0731 work (04) 380 9026 home 025 409 957 mobile
Sue Whiteman State Highway 1 McKays Crossing, Paekakariki	Support Services Duty Manager for Building	(04) 568 0744 work (04) 292 8018 home 025 411 812 mobile
Diane Lewis 386c Karori Road Wellington	National Manager Corporate Services Business & Logistics Branch	(04) 470 2514 work (04) 476 3459 home 021 342 604 mobile
Janet Campbell 100 Debbie Monahan	Support Services Manager IPONZ Manager IPONZ	(04) 560 1601 work (04) 938 7078 home (04) 560 1615 work 021 306 098 mobile
Michael Grossman Trp Met Val Asp Ser Val	Operations, BRB Glu	(09) 913 4221 025 443 702 mobile
Glas McKenzie	BRB IT System Administrator	021 532 689 mobile
Nedax Security	Mark Eden	(04) 471 2836
Anna Quanta Security	Monitoring Centre	(04) 478 1226
DX<211> 140		Phone: (04) 473 9510
Post & Waste Couriers		Phone: (04) 499 2121
ASB Cleaners	Eric Reille	(04) 564 3249

<213> Pinus radiata

<400> 187		025 454 162
Met	Pro Asn Leu Cys Leu Cys	
1	5	10
Name Address Telephone (non business)		
Arg	50	55
Diane Innes	385C Karon Rd	04 476 3459
Val	20	25
Pro Asn Leu Gly Met Cys	Karon, Wellington	021-342-634
	Arg Gly Gly Lys Ser Ile	025 243 3432
Met	25	30
Neville Harris	48 Penrose St	04 566 3460
Ser Met Ser Ser Thr Thr	Lower Hunt	021-459-158
Arg	50	55
Arg Ile Ala Gly His His	Ser Asn Leu Trp Asp Asp	025 241 8382
65	70	75
Andrew Bridgman	3 Fettes Cres	04 388 9704
Ala	85	90
Ser Leu Ser Thr Ser Tyr	Seatons Heights	021 306 722
Adam Feeley	36 Fortification Rd	04 388 2875
Asp	50	55
Lys Leu Ile Gly Glu Val	58A Asn Ile Phe Asp Leu	021 333 539
Rodney Grindley	26 Taupo Cres	04 233 9080
Glu	100	105
Asp Gly Val Phe Thr Ser	Pro Leu Ser Asp Leu His	025 433 013
Trp	115	120
Karina Bach	102 Heke Street	(04) 479 7399
Met Val Asp Ser Val Glu	Arg Leu Gly Ile Asp	025 461 868
130	135	140
	Ngaio	

<210> 188

<211> 68

<212> PRT

<213> Pinus radiata

<400> 188

Leu Gly Met Pro Arg Arg Trp Lys Phe Ala Arg Pro Ser Met Ser Leu
 1 5 10 15
 Ser Thr Val Ala Ser Asp Asp Asp Ile Gln Arg Arg Thr Gly Gly Tyr
 20 25 30
 His Ser Asn Leu Trp Asn Asp Asp Val Ile Gln Phe Leu Ser Thr Pro
 35 40 45
 Tyr Gly Glu Leu Ala Tyr Arg Glu Arg Ala Glu Arg Leu Ile Asp Glu
 50 55 60
 Val Arg Asn Ile
 65

<210> 189

<211> 99

<212> PRT

<213> Pinus radiata

<400> 189

Asp Asp Ala Val Ile Arg Arg Arg Gly Asp Tyr His Ser Asn Ile Trp
 1 5 10 15
 Asp Tyr Asp Phe Ile Gln Ser Leu Ser Ala Pro Tyr Gly Glu Pro Ser
 20 25 30
 Tyr Leu Glu Arg Ala Glu Arg Leu Ile Glu Glu Val Lys Lys Val Phe
 35 40 45
 Asn Ser Met Ser Glu Glu Asn Gly Glu Leu Ile Thr Pro Leu Asn Asp
 50 55 60
 Leu Ile Gln Arg Leu Trp Met Val Asp Ser Val Glu Arg Leu Gly Ile
 65 70 75 80
 Asp Arg His Phe Glu Asn Glu Ile Glu Ser Ala Leu Asp Tyr Val Tyr
 85 90 95
 Ser Tyr Trp

<210> 190

Ser Asn Ala Asn Gln Leu Ser Ser Met Gly Phe Ala Phe Ser Ser Gly
 20 25 30
 Ser Leu Tyr His Gln Val Met Arg Thr Lys Leu Gln Ser Met
 35 40 45
 Gly Arg Val Gly Lys Ala Tyr Ala Ser Ala Leu Ser Asp Gln Gly
 50 55 60
 Gln Tyr Ser Ser Gly Pro Thr Pro Leu Leu Asp Thr Ile Asn
 65 70 75 80
 Tyr Pro Ile His Met Lys Asn Leu Ser Ile Arg Gln Leu Lys Gln Leu
 85 90 95
 Ser Asn Gln Leu Arg Ser Asp Ile Ile Phe Gln Val Ser Arg Thr Gly
 100 105 110
 Gly His Leu Gly Ser Ser Leu Gly Val Val Gln Leu Thr Val Ala Leu
 115 120
 His Tyr Val Phe Asp Ala Pro Gln Asp Lys Ile Leu Trp Asp Val Gly
 130 135 140
 His Gln Tested on Thursday 30 September and Sunday 3 October 1999 Asp Lys Met
 145 150 155 160
 Pro Thr Lys Ser Met Arg Thr Asn Gly Leu Ser Phe Thr Lys Arg Ser
 165 170 175
 Gln Ser Gln Tyr Asp Lys Phe Gly Ala Gln His Ser Ser Thr Ser Ile
 180 185 190
 Ser Ala Gly Leu Gly Met Ala Val Gly Arg Asp Leu Lys Gly Gln Asn
 195 200 205
 Asn His Val Ile Ser Val Ile Gly Asp Gly Ala Met Thr Ala Gly Gln
 210 215 220
 Ala Phe Gln Tested on Monday 4 October 1999
 225 230
 Val Ile Leu Asn Asp Asn Lys Gln Val Ser Leu Pro Thr Ala Asn Leu
 245 250 255
 Asp Gly Pro Ile Pro Pro Val Gly Ala Leu Ser Ser Ala Leu Ser Lys
 260 265 270
 Leu Gln Ser Ser Lys Pro Leu Arg Gln Leu Arg Gln Val Ala Lys Gly
 275 280
 Val Thr Lys Gln Leu Gly Ala Pro Met His Gln Leu Ala Ala Lys Val
 290 295 300
 Asp Gln Tyr Ala Arg Gly Met Ile Ser Gly Ser Arg Ser Thr Leu Phe
 305 310 315 320
 Gln Gln Leu

<210> 193
 <211> 88
 <212> PRT
 <213> Eucalyptus grandis

<400> 193
 Gly Gly His Leu Ser Ala Ser Leu Gly Val Val Gln Leu Thr Val Ala
 1 5 10 15
 Leu His Asn Val Phe Asn Ala Pro Gln Asp Lys Ile Val Trp Asp Val
 20 25 30
 Gly His Gln Thr Tyr Pro His Lys Ile Leu Thr Gly Arg Thr Arg
 35 40 45
 Met His Thr Ile Arg Lys Thr Ser Gly Leu Ala Gly Phe Pro Lys Arg
 50 55 60
 Asp Gln Ser Val Tyr Asp Thr Phe Gly Val Gly His Ser Ser Thr Ser
 65 70 75 80
 Ile Ser Ala Gly Leu Gly Met Ala
 85

<210> 194

<211> 97
 <212> PRT
 <213> Eucalyptus grandis

<400> 194
 Pro Val Arg Glu Lys Leu Val Lys Ala Trp Arg Asn Asp Ser Glu Ile
 1 5 10 15
 Phe Ala His Tyr Gly Arg Leu Thr Thr Pro Tyr Ser Asp Glu Leu Leu
 20 25 30
 Gly Ser Lys Phe Cys Leu His Val Lys Gly Phe Glu Val Asn Thr Ala
 35 40 45
 Arg Ile Ala Asp Ser Leu Tyr Tyr Gly Cys Val Pro Val Ile Ile Ala
 50 55 60
 Asn His Tyr Asp Leu Pro Phe Ala Asp Ile Leu Asn Trp Lys Ser Phe
 65 70 75 80
 Ser Val Val Val Ala Thr Leu Asp Ile Pro Leu Leu Lys Arg Ile Leu
 85 90 95
 Lys

<210> 195
 <211> 149
 <212> PRT
 <213> Eucalyptus grandis

<400> 195
 Gly Met His Thr Ser Lys Phe Cys Leu Asn Pro Ala Gly Asp Thr Pro
 1 5 10 15
 Ser Ala Cys Arg Leu Phe Asp Ala Ile Val Ser Leu Cys Ile Pro Val
 20 25 30
 Ile Val Ser Asp Ser Ile Glu Leu Pro Phe Glu Asp Val Ile Asp Tyr
 35 40 45
 Arg Lys Ile Ala Ile Phe Val Asp Thr Ala Thr Ser Leu Lys Arg Gly
 50 55 60
 Phe Leu Val Lys Leu Leu Arg Lys Val Arg Thr Glu Lys Ile Leu Glu
 65 70 75 80
 Tyr Gln Lys Glu Leu Lys Glu Val Lys Arg Phe Phe Glu Tyr Gly Asp
 85 90 95
 Pro Asn Gly Thr Val Lys Glu Ile Trp Arg Gln Ile Ser Gln Lys Leu
 100 105 110
 Pro Leu Ile Lys Leu Met Ile Asn Arg Asp Lys Arg Ile Val Lys Arg
 115 120 125
 Asp Met Ser Glu Pro Asp Cys Ser Cys Ile Cys Ser Asn Gln Thr Gly
 130 135 140
 Val Ile Ser Thr Leu
 145

<210> 196
 <211> 196
 <212> PRT
 <213> Eucalyptus grandis

<400> 196
 Met Ser Gln Val Ser Ala Thr Pro Cys Ala Pro Pro Asn Lys Glu Thr
 1 5 10 15
 Gly His Val Ile Glu Arg Arg Ser Ala Gly Tyr His Pro Ser Val Trp
 20 25 30
 Gly Asp Tyr Phe Leu Lys Tyr Asp Ser Pro Ser Asn Ser Val Lys Phe
 35 40 45
 Lys Phe Leu Gly Arg Val Glu Gly Gln Ile Glu Glu Leu Lys Gly Glu
 50 55 60

Val Lys Lys Met Leu Ile Asp Val Val Asp Lys Pro Leu Pro Lys Leu
 65 70 75 80
 His Leu Ile Asp Gln Ile Gln Arg Leu Gly Ile Glu Tyr His Phe Glu
 85 90 95
 Arg Glu Val Asp Glu Gln Leu Glu Gln Ile His Lys Ser Tyr Ser Arg
 100 105 110
 Leu Asp His Glu Asp Phe Lys Val Asp Asp Leu His Thr Val Ala Leu
 115 120 125
 Ile Phe Arg Leu Leu Arg Gln His Gly Tyr Asn Ile Ser Ser Glu Ile
 130 135 140
 Phe Asp Lys Phe Lys Asp Ser Asn Gly Asn Phe Arg Glu Ser Leu Ile
 145 150 155 160
 Ser Asp Val Arg Gly Leu Leu Ser Leu Tyr Glu Ala Cys His Leu Arg
 165 170 175
 Cys His Gly Asp Ser Ile Leu Asp Glu Ala Leu Pro Phe Ala Thr Thr
 180 185 190
 His Leu Glu Ser
 195

<210> 197
 <211> 116
 <212> PRT
 <213> Eucalyptus grandis

<400> 197
 Met Ser Gln Val Ser Ala Thr Pro Cys Ala Pro Ser Asn Lys Gly Thr
 1 5 10 15
 Gly His Val Ile Glu Arg Arg Ser Ala Gly Tyr His Pro Ser Val Trp
 20 25 30
 Gly Asp Tyr Phe Leu Lys Tyr Asp Ser Pro Ser Asn Ser Val Lys Phe
 35 40 45
 Lys Phe Leu Gly Arg Val Glu Gly Gln Ile Glu Glu Leu Lys Gly Glu
 50 55 60
 Val Lys Lys Met Leu Thr Asp Ile Met Asp Lys Pro Leu Gln Lys Leu
 65 70 75 80
 His Leu Ile Asp Gln Ile Gln Arg Leu Gly Ile Glu Tyr His Phe Glu
 85 90 95
 Arg Glu Ile Asp Glu Gln Leu Glu Gln Ile His Lys Ser Tyr Ser Arg
 100 105 110
 Leu Asp His Glu
 115

<210> 198
 <211> 102
 <212> PRT
 <213> Eucalyptus grandis

<400> 198
 Met Ser Leu Pro Ile Ser Arg Val Pro Ser Ser Ser Pro Ala Glu Lys
 1 5 10 15
 Thr Ser Leu Val Pro Glu Gly Gly Ser Ala Ile Phe His Pro Thr Ile
 20 25 30
 Trp Ala Asp Tyr Phe Leu Lys His Ala Ser Asn Ser Asn Ser Thr Ser
 35 40 45
 Ser Asp Gly Val Val Glu Glu His Ile Glu Arg Leu Lys Gly Glu Val
 50 55 60
 Arg Lys Met Leu Met Gly Ala Met Asp Lys Pro Ser Gln Lys Leu Asn
 65 70 75 80
 Leu Ile Asp Gln Ile Gln Arg Leu Gly Phe Ala Tyr His Phe Glu His
 85 90 95
 Glu Ile Asp Glu Gln Leu

100

<210> 199
 <211> 169
 <212> PRT
 <213> Eucalyptus grandis

<400> 199

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Thr Ser Phe Leu Pro Ser Ser Ile His His Asn Gln Pro Ser Leu Leu
 1          5          10          15
Phe Phe Arg His Leu Cys Ser Ser Ser Ala Ala Thr Ser Ser Thr
      20          25          30
Ser Ser Gly Ala Gln Phe Val Thr Cys Thr Leu Lys Ile Glu Ala Gln
      35          40          45
Glu Ile Gly Arg Arg Ser Ala Asn Trp Gln Pro Asn Val Phe Asp Tyr
      50          55          60
Asp Phe Leu Gln Ser Leu Asn Val Asp Tyr Thr Glu Asp Lys Tyr Ser
      65          70          75          80
Glu Glu Ala Gln Arg Leu Lys Lys Glu Val Lys Gly Leu Phe Asp Lys
      85          90          95
Lys Met Asn Ser Val Ala Lys Leu Glu Phe Ile Asp Val Val Gln Arg
      100          105          110
Leu Gly Leu Gly Tyr Gln Phe Glu Thr Glu Ile Lys Asn Ala Leu Ser
      115          120          125
Ser Ile Tyr Asn Asn Ala Glu Asp Ala Gln Leu Leu Asp Asp Leu Tyr
      130          135          140
Ala Val Ser Leu Arg Phe Arg Leu Leu Arg Gln His Gly Phe Asn Ile
      145          150          155          160
Ser Gln Asp Ala Phe Gln Arg Phe Met
      165

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<210> 200
 <211> 132
 <212> PRT
 <213> Eucalyptus grandis

<400> 200

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Ser Ile Arg Pro Asn Gln Pro Ser Leu Ser Leu Phe Ser Arg Pro Arg
 1          5          10          15
Ser Ser Phe Ser Ser Pro Ser Ala Val Ser Ser Gly Thr Arg Phe Ala
      20          25          30
Lys Cys Ala Leu Thr Ile Glu Asp Glu Asp Thr Ala Arg Arg Ser Ala
      35          40          45
Asn Trp Lys Pro Ser Val Trp Asp Tyr Gly Phe Val Gln Ser Leu Asn
      50          55          60
Thr Asp Phe Pro Val Asp Lys Tyr Thr Glu Gln Val Gln Arg Leu Lys
      65          70          75          80
Glu Glu Val Lys Gly Leu Phe His Arg Glu Met Asn Gln Val Ala Lys
      85          90          95
Leu Glu Phe Ile Asp Val Val Gln Arg Leu Gly Leu Gly Tyr His Phe
      100          105          110
Glu Thr Glu Ile Asn Asn Ser Leu Ser Ser Ile Tyr Asn Asn Thr Glu
      115          120          125
Asp Val Gln Leu
      130

```

<210> 201
 <211> 116
 <212> PRT
 <213> Pinus radiata

<400> 201

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Met Ala Ser Val Ser Val Lys Ala Gly Ala Thr Ser Thr Val Ser Cys
 1           5           10           15
Gly Leu Ala Ser Asn Asn Leu Ile Arg Arg Thr Ala Asn Pro His Pro
          20           25           30
Asn Val Trp Asp Tyr Asp Phe Val His Ser Leu Lys Ser Pro Tyr Asn
          35           40           45
Asp Ser Ser Tyr Thr Glu Arg Ala Glu Thr Leu Ile Gly Gln Leu Lys
          50           55           60
Val Met Leu Ser Ala Ala Ile Gly Gly Gly Glu Ser Met Ile Thr Pro
          65           70           75           80
Ser Ala Tyr Asp Thr Ala Trp Val Ala Arg Val Pro Ser Ile Asp Gly
          85           90           95
Ser Ala Cys Pro Gln Phe Pro Gln Thr Val Glu Trp Ile Leu Lys Asn
          100          105          110
Gln Leu Lys Asp
          115

```

<210> 202

<211> 121

<212> PRT

<213> Pinus radiata

<400> 202

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Ala Ile Leu Ser Tyr Pro Pro Glu Ile Leu Ala Leu Pro Ser Pro Ser
 1           5           10           15
Phe Leu Tyr Ile Ser Ser Leu Ile Pro Met Ala Ser Val Val Asp Gln
          20           25           30
Ala Glu Leu Cys Ser Lys Ser Val Ser Met Ser Ser Pro Gly Val Gln
          35           40           45
Arg Arg Thr Gly Asp Tyr His Ser Asn Leu Trp Asp Asp Glu Phe Ile
          50           55           60
Gln Ser Leu Ser Thr Pro Tyr Gly Ala Pro Ser Tyr Arg Glu Arg Ala
          65           70           75           80
Asp Arg Leu Val Gly Glu Val Lys Glu Met Phe Asn Ser Leu Thr Val
          85           90           95
Leu Thr Pro His Asn Asp Leu Leu Glu Gln Leu Trp Met Val Asp Ser
          100          105          110
Val Glu Arg Leu Gly Ile Asp Arg His
          115          120

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<210> 203

<211> 259

<212> PRT

<213> Pinus radiata

<400> 203

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Asn Ile Gly Pro Ser Phe Leu Ser Ile Ser Ser Leu Val Arg Met Ala
 1           5           10           15
Ser Val Val Asp Gln Ala Glu Leu Cys Ser Lys Ser Val Ser Met Ser
          20           25           30
Ser Pro Gly Val Gln Arg Arg Thr Gly Asp Tyr His Ser Asn Leu Trp
          35           40           45
Asp Asp Asp Phe Ile Gln Ser Leu Ser Thr Pro Tyr Gly Ala Pro Ser
          50           55           60
Tyr Arg Glu Arg Ala Asp Arg Leu Val Gly Glu Val Lys Glu Met Phe
          65           70           75           80
Asn Ser Leu Thr Leu Leu Thr Pro Leu Asn Asp Leu Leu Gln Arg Leu
          85           90           95
Trp Met Val Asp Thr Val Glu Arg Leu Glu Ile Asp Arg His Phe Arg
          100          105          110

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Asn Glu Ile Lys Ser Ala Leu Asp Tyr Val Tyr Ser Tyr Trp Ser Glu
 115 120 125
 Lys Gly Ile Gly Cys Gly Arg Glu Ser Val Val Thr Asp Leu Asn Ser
 130 135 140
 Thr Ala Leu Gly Phe Arg Thr Leu Arg Leu His Gly Phe Pro Val Ser
 145 150 155 160
 Ser Asp Val Leu Glu Val Phe Lys Asp Gln Asn Gly Lys Phe Ala Gly
 165 170 175
 Cys Ser Ala Asn Ala Glu Thr Glu Ala Glu Met Arg Asp Ile Leu Asn
 180 185 190
 Leu Phe Arg Ala Ser Leu Val Ala Phe Pro Gly Glu Lys Val Met Glu
 195 200 205
 Glu Ala Gln Thr Phe Cys Thr Ser Tyr Leu Gln Glu Ala Leu Lys Thr
 210 215 220
 Val Pro Ile Ser Asn Asp Ser Leu Ser Arg Glu Ile Glu Tyr Val Ile
 225 230 235 240
 Glu Tyr Gly Trp Leu Thr Asn Phe Ser Glu Ile Gly Ser Lys Glu Leu
 245 250 255
 His Arg Arg

<210> 204
 <211> 344
 <212> PRT
 <213> Pinus radiata

<400> 204
 Ile Asp Val Phe Gly Glu Asp Thr Thr Phe Glu Thr Pro Tyr Leu Ile
 1 5 10 15
 Arg Glu Lys Leu Leu Glu Leu Ala Lys Leu Glu Phe Asn Ile Phe His
 20 25 30
 Ser Leu Val Lys Arg Glu Leu Gln Ser Leu Leu Arg Trp Trp Lys Asp
 35 40 45
 Tyr Gly Phe Pro Glu Ile Thr Phe Ser Arg His Arg His Val Glu Tyr
 50 55 60
 Tyr Thr Leu Ala Ala Cys Ile Ala Asn Asp Pro Lys His Ser Ala Phe
 65 70 75 80
 Arg Leu Gly Phe Gly Lys Ile Ser His Met Ile Thr Ile Leu Asp Asp
 85 90 95
 Ile Tyr Asp Thr Phe Gly Thr Met Glu Glu Leu Glu Leu Leu Thr Ala
 100 105 110
 Ala Phe Lys Arg Trp Asp Pro Ser Ser Ile Glu Cys Leu Pro Asp Tyr
 115 120 125
 Met Lys Gly Val Tyr Met Ala Val Tyr Asp Asn Ile Asn Glu Met Ala
 130 135 140
 Arg Glu Ala Gln Lys Ile Gln Gly Trp Asp Thr Val Ser Tyr Ala Arg
 145 150 155 160
 Lys Ser Trp Glu Ala Phe Ile Gly Ala Tyr Ile Gln Glu Ala Lys Trp
 165 170 175
 Ile Ser Ser Gly Tyr Leu Pro Thr Phe Asp Glu Tyr Leu Glu Asn Gly
 180 185 190
 Lys Val Ser Phe Gly Ser Arg Ile Thr Thr Leu Glu Pro Met Leu Thr
 195 200 205
 Leu Gly Phe Pro Leu Pro Pro Arg Ile Leu Gln Glu Ile Asp Phe Pro
 210 215 220
 Pro Lys Phe Asn Asp Leu Ile Cys Ala Ile Leu Arg Leu Lys Gly Asp
 225 230 235 240
 Thr Gln Cys Tyr Lys Ala Asp Arg Ala Arg Gly Glu Glu Ala Ser Ala
 245 250 255
 Val Ser Cys Tyr Met Lys Asp His Pro Gly Ile Thr Glu Glu Asp Ala
 260 265 270

Val Asn Gln Val Asn Ala Met Val Asp Asn Leu Thr Lys Glu Leu Asn
 275 280 285
 Trp Glu Leu Leu Arg Pro Asp Ser Gly Val Pro Ile Ser Tyr Lys Lys
 290 295 300
 Val Ala Phe Asp Ile Cys Arg Val Phe His Tyr Gly Tyr Lys Tyr Arg
 305 310 315 320
 Asp Gly Phe Ser Val Ala Ser Ile Glu Ile Lys Asn Leu Val Thr Arg
 325 330 335
 Thr Val Val Glu Thr Val Pro Leu
 340

<210> 205
 <211> 462
 <212> PRT
 <213> Pinus radiata

<400> 205
 Arg Asp Ser Ala Phe Thr Asp Leu Asn Thr Thr Ala Leu Gly Phe Arg
 1 5 10 15
 Ile Phe Arg Leu His Gly Tyr Thr Val Ser Ser Asp Ala Phe Glu His
 20 25 30
 Phe Lys Asp Gln Met Gly Gln Phe Ser Ala Ser Ala Asn Asp Thr Glu
 35 40 45
 Leu Gln Ile Arg Ser Val Phe Asn Leu Phe Arg Ala Ser Leu Ile Ala
 50 55 60
 Phe Pro Glu Glu Lys Val Leu Glu Glu Ala Glu Asn Phe Ala Ala Ala
 65 70 75 80
 Tyr Leu Lys Ala Ala Leu Gln Thr Leu Pro Val Ser Gly Leu Ser Arg
 85 90 95
 Glu Ile Gln Tyr Val Phe Asp Tyr Arg Trp His Ser Asn Leu Pro Arg
 100 105 110
 Leu Glu Ala Arg Ser Tyr Val Asp Ile Leu Ala Asp Asn Thr Ile Ser
 115 120 125
 Gly Thr Pro Asp Ala Asn Thr Lys Lys Leu Leu Glu Leu Ala Lys Leu
 130 135 140
 Glu Phe Asn Ile Phe His Ser Leu Gln Gln Lys Glu Leu Gln Cys Leu
 145 150 155 160
 Trp Arg Trp Trp Lys Glu Trp Gly Cys Pro Glu Leu Thr Phe Val Arg
 165 170 175
 His Arg Tyr Val Glu Phe Tyr Thr Leu Val Ser Gly Thr Asp Met Val
 180 185 190
 Pro Glu His Ala Ala Phe Arg Leu Ser Phe Val Lys Thr Cys His Leu
 195 200 205
 Ile Thr Ile Leu Asp Asp Met Tyr Asp Thr Phe Gly Thr Ile Asp Glu
 210 215 220
 Leu Arg Leu Phe Thr Ala Ala Val Lys Arg Trp Asp Pro Ser Ala Thr
 225 230 235 240
 Glu Cys Leu Pro Glu Tyr Met Lys Gly Val Tyr Arg Val Leu Tyr Glu
 245 250 255
 Thr Val Asn Glu Met Ala Lys Glu Ala Gln Lys Ser Gln Gly Arg Asp
 260 265 270
 Thr Leu Gly Tyr Val Arg Gln Ala Leu Glu Asp Tyr Ile Gly Ser Tyr
 275 280 285
 Leu Lys Glu Ala Glu Trp Ile Ala Thr Gly Tyr Val Pro Thr Phe Gln
 290 295 300
 Glu Tyr Phe Glu Asn Gly Lys Leu Ser Ser Gly His Arg Ile Ala Thr
 305 310 315 320
 Leu Gln Pro Ile Leu Thr Leu Ser Ile Pro Phe Pro His His Ile Leu
 325 330 335
 Gln Glu Ile Asp Phe Pro Ser Lys Phe Asn Asp Tyr Ala Cys Ser Ile
 340 345 350

Leu Arg Leu Arg Gly Asp Thr Arg Cys Tyr Lys Ala Asp Ser Ala Arg
 355 360 365
 Gly Glu Glu Ala Ser Cys Ile Ser Cys Tyr Met Lys Glu Asn Pro Gly
 370 375 380
 Ser Thr Gln Glu Asp Ala Leu His His Ile Asn Gly Met Ile Glu Asp
 385 390 395 400
 Met Ile Lys Lys Leu Asn Trp Glu Phe Leu Lys Pro Asp Asn Asn Ala
 405 410 415
 Pro Ile Ser Ser Lys Lys Asn Ala Phe Asn Ile Ser Arg Gly Leu His
 420 425 430
 His Phe Tyr Asn Tyr Arg Asp Gly Tyr Ser Val Ala Ser Asn Glu Thr
 435 440 445
 Lys Asp Leu Val Ile Lys Thr Val Leu Glu Pro Val Leu Met
 450 455 460

<210> 206
 <211> 100
 <212> PRT
 <213> Eucalyptus grandis

<400> 206
 Gly Ser Gln Leu Trp Asp Thr Ala Phe Ala Thr Gln Ala Ile Ile Ser
 1 5 10 15
 Thr Asn Leu Ile Glu Glu Phe Gly Ser Thr Leu Gln Lys Ala His Thr
 20 25 30
 Tyr Ile Lys Asn Ser Gln Val Leu Glu Asp Cys Pro Gly Asp Leu Asn
 35 40 45
 Phe Trp Tyr Arg His Ile Ser Lys Gly Ala Trp Pro Phe Ser Thr Ala
 50 55 60
 Asp His Gly Trp Pro Ile Ser Asp Cys Thr Ala Glu Gly Leu Lys Ala
 65 70 75 80
 Ala Leu Val Leu Ser Lys Ile Pro Leu Glu Ile Val Gly Gln Pro Phe
 85 90 95
 Arg Ser Tyr Gly
 100

<210> 207
 <211> 89
 <212> PRT
 <213> Eucalyptus grandis

<400> 207
 Met Trp Lys Leu Lys Val Ala Glu Gly Ala Asn Pro Trp Leu Arg Ser
 1 5 10 15
 Leu Asn Asn His Val Gly Arg Gln Ile Trp Glu Phe Asp Pro Asn Cys
 20 25 30
 Gly Ser Pro Glu Glu Ile Gln Glu Ile Glu Glu Ala Arg Ala Asn Phe
 35 40 45
 Leu Lys His Arg Phe Glu Lys Lys His Ser Ser Asp Leu Met Met Arg
 50 55 60
 Ile Gln Phe Ser Lys Glu Asn Thr Gly Arg Val Val Leu Pro Pro Val
 65 70 75 80
 Lys Val Lys Asp Leu Asp Glu Ile Thr
 85

<210> 208
 <211> 198
 <212> PRT
 <213> Eucalyptus grandis

<400> 208

Val Thr His Met Leu Arg Arg Ala Ile Ser Phe His Ser Thr Leu Gln
 1 5 10 15
 Ala His Asp Gly His Trp Pro Gly Asp Tyr Gly Gly Pro Met Phe Leu
 20 25 30
 Met Pro Gly Leu Val Ile Ala Leu Ser Ile Thr Gly Ala Leu Asn Ala
 35 40 45
 Val Leu Ser Glu Gln His Lys Gln Glu Met Cys Arg Tyr Leu Tyr Asn
 50 55 60
 His Gln Asn Lys Asp Gly Gly Trp Gly Leu His Ile Glu Gly Pro Ser
 65 70 75 80
 Thr Met Phe Gly Ser Val Leu Asn Tyr Val Thr Leu Arg Leu Leu Gly
 85 90 95
 Glu Ala Ala Asn Asp Gly Gln Gly Ala Met Glu Lys Ala Arg Lys Trp
 100 105 110
 Ile Leu Asp His Gly Ser Ala Thr Ala Ile Thr Ser Trp Gly Lys Met
 115 120 125
 Trp Leu Ser Val Leu Gly Ala Phe Glu Trp Ser Gly Asn Asn Pro Leu
 130 135 140
 Pro Pro Glu Ile Trp Leu Leu Pro Tyr Met Leu Pro Ile His Pro Gly
 145 150 155 160
 Arg Met Trp Cys His Cys Arg Met Val Tyr Leu Pro Met Ser Tyr Leu
 165 170 175
 Tyr Gly Lys Arg Phe Val Ser Pro Ile Thr Pro Thr Val Phe Val Leu
 180 185 190
 Glu Lys Arg Asn Phe Met
 195

<210> 209

<211> 78

<212> PRT

<213> Eucalyptus grandis

<400> 209

Met Trp Lys Leu Lys Ile Ala Glu Gly Gly Pro Trp Leu Thr Ser Val
 1 5 10 15
 Asn Asn His Val Gly Arg Gln His Trp Glu Phe Asp Pro Asp Ala Gly
 20 25 30
 Thr Pro Glu Glu Arg Ala Glu Val Glu Arg Val Arg Asp Glu Phe Thr
 35 40 45
 Arg Asn Arg Phe Arg Ile Lys Gln Ser Ala Asp Leu Leu Met Arg Met
 50 55 60
 Gln Leu Thr Lys Glu Asn Pro Ser Gly Pro Ile His Arg Arg
 65 70 75

<210> 210

<211> 97

<212> PRT

<213> Eucalyptus grandis

<400> 210

Tyr Val Trp Val Gly Glu Asp Gly Ile Lys Met Gln Ser Phe Gly Ser
 1 5 10 15
 Gln Ile Trp Asp Cys Gly Leu Ser Leu Gln Ala Leu Leu Ala Ser Asp
 20 25 30
 Leu Ile Asp Glu Ile Gly Pro Val Leu Lys Lys Gly His Glu Phe Leu
 35 40 45
 Lys Glu Ser Gln Ile Asp Arg Asn Pro Ser Gly Asp Leu Lys Lys Met
 50 55 60
 Tyr Arg His Ile Ser Lys Gly Ala Trp Ala Phe Ser Asp Lys Asp His
 65 70 75 80
 Gly Trp Gln Val Ser Asp Cys Thr Ala Glu Ser Met Lys Cys Cys Leu

Val 85 90 95

<210> 211
 <211> 158
 <212> PRT
 <213> Eucalyptus grandis

<400> 211

Met	Asp	Thr	Asp	Asn	Lys	Leu	Phe	Asn	Val	Gly	Val	Leu	Leu	Val	Ala
1				5					10					15	
Thr	Leu	Val	Val	Ala	Lys	Leu	Ile	Ser	Ala	Leu	Leu	Ile	Pro	Arg	Ser
			20					25					30		
Gly	Lys	Arg	Leu	Pro	Pro	Val	Val	Arg	Thr	Trp	Pro	Val	Val	Gly	Gly
	35					40					45				
Leu	Leu	Arg	Phe	Leu	Lys	Gly	Pro	Met	Val	Met	Leu	Arg	Glu	Glu	Tyr
	50					55				60					
Pro	Lys	Leu	Gly	Ser	Val	Phe	Thr	Leu	Asn	Leu	Leu	Asn	Lys	Lys	Ile
65					70					75					80
Thr	Phe	Phe	Ile	Gly	Pro	Glu	Val	Ser	Ala	His	Phe	Phe	Lys	Ala	Ser
			85					90						95	
Glu	Ser	Asp	Leu	Ser	Gln	Gln	Glu	Val	Tyr	Gln	Phe	Asn	Val	Pro	Thr
			100					105					110		
Phe	Gly	Pro	Gly	Val	Val	Phe	Asp	Val	Asp	Tyr	Thr	Ile	Arg	Gln	Glu
	115					120						125			
Gln	Phe	Arg	Phe	Phe	Thr	Glu	Ala	Leu	Arg	Ile	Asn	Lys	Leu	Lys	Gly
	130					135					140				
Tyr	Val	Asn	Gln	Met	Val	Met	Glu	Ala	Glu	Asp	Tyr	Phe	Ser		
145					150					155					

<210> 212
 <211> 131
 <212> PRT
 <213> Eucalyptus grandis

<400> 212

Met	Asp	Thr	Asp	Asn	Lys	Leu	Phe	Asn	Val	Gly	Val	Leu	Leu	Val	Ala
1				5					10					15	
Thr	Leu	Val	Val	Ala	Lys	Leu	Ile	Ser	Ala	Leu	Leu	Ile	Pro	Arg	Ser
			20					25					30		
Gly	Lys	Arg	Leu	Pro	Pro	Val	Val	Arg	Thr	Trp	Pro	Val	Val	Gly	Gly
	35					40					45				
Leu	Leu	Arg	Phe	Leu	Lys	Gly	Pro	Met	Val	Met	Leu	Arg	Glu	Glu	Tyr
	50					55				60					
Pro	Lys	Leu	Gly	Ser	Val	Phe	Thr	Leu	Asn	Leu	Leu	Asn	Lys	Lys	Ile
65					70					75					80
Thr	Phe	Phe	Ile	Gly	Pro	Glu	Val	Ser	Ala	His	Phe	Phe	Lys	Ala	Ser
			85					90						95	
Glu	Ser	Asp	Leu	Ser	Gln	Gln	Glu	Val	Tyr	Gln	Phe	Asn	Val	Pro	Thr
			100					105					110		
Phe	Gly	Pro	Gly	Val	Val	Phe	Asp	Val	Asp	Tyr	Thr	Ile	Arg	Gln	Glu
	115					120						125			
Gln	Phe	Arg													
130															

<210> 213
 <211> 112
 <212> PRT
 <213> Eucalyptus grandis

<400> 213

Met Asp Thr Asp Asn Lys Leu Phe Asn Val Gly Val Leu Leu Val Ala
 1 5 10 15
 Thr Leu Val Val Ala Lys Leu Ile Ser Ala Ser Ile Pro Arg Ser Gly
 20 25 30
 Lys Arg Leu Pro Pro Val Val Arg Thr Trp Pro Val Val Gly Gly Leu
 35 40 45
 Leu Arg Phe Leu Lys Gly Pro Met Val Met Leu Arg Glu Glu Tyr Pro
 50 55 60
 Lys Leu Gly Ser Val Phe Thr Leu Asn Leu Leu Asn Lys Lys Ile Thr
 65 70 75 80
 Phe Phe Ile Gly Pro Glu Val Ser Ala His Phe Phe Lys Ala Ser Glu
 85 90 95
 Ser Asp Leu Ser Gln Gln Glu Val Tyr Gln Phe Asn Val Pro Thr Phe
 100 105 110

<210> 214

<211> 152

<212> PRT

<213> Eucalyptus grandis

<400> 214

Phe Leu Lys Gly Pro Met Val Met Leu Arg Glu Glu Tyr Pro Lys Leu
 1 5 10 15
 Gly Ser Val Phe Thr Leu Asn Leu Leu Asn Lys Lys Ile Thr Phe Phe
 20 25 30
 Ile Gly Pro Glu Val Ser Ala His Phe Phe Lys Ala Ser Glu Ser Asp
 35 40 45
 Leu Ser Gln Gln Glu Val Tyr Gln Phe Asn Val Pro Thr Phe Gly Pro
 50 55 60
 Gly Val Val Phe Asp Val Asp Tyr Thr Ile Arg Glu Glu Gln Phe Arg
 65 70 75 80
 Phe Phe Thr Glu Ala Leu Arg Ile Asn Lys Leu Lys Gly Tyr Val Asn
 85 90 95
 Gln Met Val Met Glu Ala Glu Asp Tyr Phe Ser Lys Trp Gly Asp Ser
 100 105 110
 Gly Glu Val Asp Leu Lys Tyr Glu Leu Glu His Leu Thr Ile Leu Thr
 115 120 125
 Ala Ser Arg Cys Leu Leu Gly Arg Glu Val Arg Glu Lys Leu Phe Asp
 130 135 140
 Asp Val Ser Ala Leu Phe His Asp
 145 150

<210> 215

<211> 147

<212> PRT

<213> Eucalyptus grandis

<400> 215

Phe Asp Asp Val Ser Ala Leu Phe His Asp Leu Asp Asn Gly Met Leu
 1 5 10 15
 Pro Ile Ser Val Ile Phe Pro Tyr Leu Pro Ile Pro Ala His His Arg
 20 25 30
 Arg Asp Lys Ala Arg Lys Lys Leu Ser Glu Ile Phe Ala Asn Ile Ile
 35 40 45
 Ser Ser Arg Lys Cys Ala Gly Lys Ser Glu Glu Asp Met Leu Gln Cys
 50 55 60
 Phe Ile Asp Ser Lys Tyr Lys Asn Gly Arg Pro Thr Thr Glu Ala Glu
 65 70 75 80
 Val Thr Gly Leu Leu Ile Ala Ala Leu Phe Ala Gly Gln His Thr Ser
 85 90 95

Ser Ile Thr Ser Val Trp Thr Gly Ala Tyr Leu Leu Thr Asn Lys Lys
 100 105 110
 Tyr Leu Ser Ala Val Ser Asn Glu Gln Lys His Leu Met Glu Lys His
 115 120 125
 Gly Asn Asn Val Asp His Asp Val Leu Ser Glu Met Asp Val Leu Tyr
 130 135 140
 Arg Ser Ile
 145

<210> 216
 <211> 129
 <212> PRT
 <213> Eucalyptus grandis

<400> 216
 Tyr Leu Leu Thr Asn Lys Lys Tyr Leu Ser Ala Val Ser Asn Glu Gln
 1 5 10 15
 Lys His Leu Met Glu Lys His Gly Asn Asn Val Asp His Asp Val Leu
 20 25 30
 Ser Glu Met Asp Val Leu Tyr Arg Ser Ile Lys Glu Ala Leu Arg Leu
 35 40 45
 His Pro Pro Leu Ile Met Leu Leu Arg Ser Ser His Ser Asp Phe Ser
 50 55 60
 Val Lys Thr Arg Asp Gly Lys Glu Tyr Glu Val Gly Glu Val Ser Val
 65 70 75 80
 Leu Pro Trp Thr Leu Glu Ala Arg Lys Gly Val Gly Lys Ala Phe Ile
 85 90 95
 Thr Ala Phe Arg Ser Gly Ala Val Met Gly Phe Leu Leu Ala Ala Asn
 100 105 110
 Gly Leu Leu Val Leu Tyr Ile Ala Ile Asn Leu Phe Lys Ile Tyr Leu
 115 120 125
 Trp

<210> 217
 <211> 118
 <212> PRT
 <213> Eucalyptus grandis

<400> 217
 Val Val Phe Asp Val Asp Tyr Thr Ile Arg Gln Glu Gln Phe Arg Phe
 1 5 10 15
 Phe Thr Glu Ala Leu Arg Ile Asn Lys Leu Lys Gly Tyr Val Asn Gln
 20 25 30
 Met Val Met Glu Ala Glu Asp Tyr Phe Ser Lys Trp Gly Asp Ser Gly
 35 40 45
 Glu Val Asp Leu Lys Tyr Glu Leu Glu His Leu Thr Ile Leu Thr Ala
 50 55 60
 Ser Arg Cys Leu Leu Gly Arg Glu Val Arg Glu Lys Leu Phe Asp Asp
 65 70 75 80
 Val Ser Ala Leu Phe His Asp Leu Asp Asn Gly Met Leu Pro Ile Ser
 85 90 95
 Val Ile Phe Pro Tyr Leu Pro Ile Pro Ala His His Arg Arg Asp Lys
 100 105 110
 Ala Arg Lys Lys Leu Ala
 115

<210> 218
 <211> 146
 <212> PRT
 <213> Eucalyptus grandis

<400> 218
 Ser Val Arg Arg Arg Ala Leu Glu Met Thr Thr Gly Arg Cys Leu Asp
 1 5 10 15
 Gly Leu Pro Leu Asp Gly Phe Asp Tyr Gly Ser Ile Leu Gly Gln Cys
 20 25 30
 Cys Glu Leu Leu Pro Ile Gly Tyr Val Gln Ile Pro Val Gly Val Ala Gly
 35 40 45
 Pro Leu Leu Leu Asp Gly Ile Glu Asn Met Val Pro Met Ala Thr Thr
 50 55 60
 Glu Gly Cys Leu Val Ala Ser Thr Asn Arg Gly Cys Lys Ala Ile His
 65 70 75 80
 Met Ser Gly Gly Ala Thr Ser Val Leu Leu Arg Asp Gly Met Thr Arg
 85 90 95
 Ala Pro Val Val Arg Phe Pro Thr Ala Arg Arg Ala Ala Gln Leu Lys
 100 105 110
 Phe Tyr Leu Glu Ala Pro Ile Thr Thr Lys Ala Cys Leu Ser Ser Ser
 115 120 125
 Thr Ala Pro Ser Lys Val Cys Gln Ala Cys Lys Gly Ile Gln Val Pro
 130 135 140
 Pro Ile
 145

<210> 219
 <211> 328
 <212> PRT
 <213> Eucalyptus grandis

<400> 219
 Val Ala Ser Tyr Ser Leu Glu Ser Ala Leu Gly Gly Asp Cys Arg Arg
 1 5 10 15
 Ala Ala Leu Val Arg Arg Arg Ala Leu Glu Ile Arg Thr Gly Arg Cys
 20 25 30
 Leu Asp Gly Leu Pro Leu Asp Gly Phe Asp Tyr Gly Ser Ile Leu Gly
 35 40 45
 Gln Cys Cys Glu Leu Pro Val Gly Tyr Val Gln Ile Pro Val Gly Val
 50 55 60
 Val Gly Pro Leu Leu Leu Asp Gly Leu Glu Asn Met Val Pro Met Ala
 65 70 75 80
 Thr Thr Glu Gly Cys Leu Val Ala Ser Ala Asn Arg Gly Cys Lys Ala
 85 90 95
 Ile His Met Ser Gly Gly Ala Thr Ser Val Leu Leu Arg Asp Gly Met
 100 105 110
 Thr Arg Ala Pro Val Val Arg Phe Pro Thr Ala Glu Arg Ala Ala His
 115 120 125
 Leu Lys Ser Tyr Leu Glu His Pro Lys Asn Phe Asp Ser Leu Ser Leu
 130 135 140
 Ile Phe Asn Ser Thr Ser Arg Phe Ala Arg Leu Gln Thr Ile Lys Cys
 145 150 155 160
 Ala Ile Ala Gly Arg Asn Leu Tyr Ile Arg Phe Ser Cys Phe Thr Gly
 165 170 175
 Asp Ala Met Gly Met Asn Met Val Ser Lys Gly Val Gln Asn Val Leu
 180 185 190
 Asp Phe Leu Gln Asn Glu Asn Pro Asp Met Asp Val Ile Ala Val Ser
 195 200 205
 Gly Asn Phe Cys Ala Asp Lys Lys Pro Thr Ala Val Asn Trp Ile Glu
 210 215 220
 Gly Arg Gly Lys Ser Val Val Cys Glu Ala Ile Ile Thr Glu Ala Val
 225 230 235 240
 Val Ser Lys Val Leu Lys Thr Thr Val Pro Ala Leu Leu Glu Leu Asn
 245 250 255

Met Leu Lys Asn Leu Thr Gly Ser Ala Leu Ala Gly Ala Met Gly Gly
 260 265 270
 Phe Asn Ala His Ala Ser Asn Ile Val Ser Ala Val Phe Ile Ala Thr
 275 280 285
 Gly Gln Asp Pro Ala Gln Asn Ile Glu Ser Ser His Cys Ile Thr Met
 290 295 300
 Met Glu Ala Ser Asn Asp Gly Lys Asp Leu His Val Ser Val Thr Met
 305 310 315 320
 Pro Cys Ile Glu Val Gly Asn Ser
 325

<210> 220

<211> 175

<212> PRT

<213> Eucalyptus grandis

<400> 220

Leu Gly Gly Asp Cys Arg Arg Ala Ala Ser Val Arg Arg Arg Ala Leu
 1 5 10 15
 Glu Met Thr Thr Gly Arg Cys Leu Asp Gly Leu Pro Leu Asp Gly Phe
 20 25 30
 Asp Tyr Gly Ser Ile Leu Gly Gln Cys Cys Glu Leu Pro Val Gly Tyr
 35 40 45
 Val Gln Ile Pro Val Gly Val Ala Gly Pro Leu Leu Asp Gly Phe
 50 55 60
 Glu Ile Met Val Pro Met Ala Thr Thr Glu Gly Cys Leu Val Ala Ser
 65 70 75 80
 Thr Asn Arg Gly Cys Lys Ala Ile His Met Ser Gly Gly Ala Thr Ser
 85 90 95
 Val Leu Leu Arg Asp Gly Met Thr Arg Ala Pro Val Val Arg Phe Ser
 100 105 110
 Thr Ala Arg Arg Ala Ala Gln Leu Lys Phe Tyr Leu Glu His Pro Asn
 115 120 125
 Asn Tyr Lys Ser Leu Ser Leu Ile Phe Asn Ser Thr Ser Arg Phe Ala
 130 135 140
 Arg Leu Gln Gly Ile Lys Cys Ala Ile Ala Gly Arg Asn Leu Tyr Met
 145 150 155 160
 Arg Phe Cys Cys Ser Thr Gly Asp Ala Met Gly Asp Glu Tyr Gly
 165 170 175

<210> 221

<211> 220

<212> PRT

<213> Pinus radiata

<400> 221

Met Glu Ser Cys Gly Ser Gly Ile Ser Gly Thr Gly Lys Lys Met Lys
 1 5 10 15
 Asn Ser Arg Thr Leu Ala Ser Asp Ala Leu Pro Leu Pro Val Gly Leu
 20 25 30
 Thr Asn Lys Val Phe Phe Ile Leu Phe Phe Thr Ala Ser Tyr Phe Leu
 35 40 45
 Met Arg Arg Trp Arg Glu Lys Ile Arg Thr Ser Thr Pro Leu His Val
 50 55 60
 Leu Ser Leu Gly Glu Leu Val Ala Ile Val Ala Gln Leu Ala Ser Phe
 65 70 75 80
 Ile Tyr Leu Leu Gly Phe Phe Gly Ile Asp Tyr Val Gln Asn Phe Ile
 85 90 95
 Thr Gly Gly Asn Asp Asp Asp Ala Arg Glu Asp Asp Lys Leu Arg
 100 105 110
 Ser Pro Val Pro Lys Glu Ala Val Ala Ile Arg Pro Ser Ala Pro Gln

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      115      120      125
Val Gln Leu Asn Gly Ile Ser Leu Gly Asp Asn Lys Asp Asp Asp Ile
      130      135      140
Ala Ala Ala Val Cys Asn Gly Thr Val Ala Ser Tyr Ser Leu Glu Ser
145      150      155      160
Ser Leu Gly Asp Cys Met Arg Ser Ala Arg Val Arg Arg Arg Ser Leu
      165      170      175
Glu Met Met Thr Gly Arg Ser Leu Asp Gly Leu Pro Leu Glu Gly Phe
      180      185      190
Asp Tyr Gly Ser Ile Leu Gly Gln Cys Cys Glu Leu Pro Val Gly Tyr
      195      200      205
Val Gln Ile Pro Val Gly Val Ala Gly Pro Leu Leu
      210      215      220

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<210> 222
 <211> 91
 <212> PRT
 <213> Pinus radiata

```

      <400> 222
Asp Leu His Ile Ser Val Thr Met Pro Cys Ile Glu Val Gly Thr Val
 1      5      10      15
Gly Gly Gly Thr Gln Leu Ala Ser Gln Ser Ala Cys Leu Asn Leu Ile
      20      25      30
Gly Val Lys Gly Ala Asn Val Gln Ser Pro Gly Ala Asn Ala Arg Leu
      35      40      45
Leu Ala Arg Ile Val Ala Gly Ala Val Leu Ala Gly Glu Leu Ser Leu
      50      55      60
Met Ser Ala Leu Ala Ala Gly Gln Leu Val Lys Ser His Met Lys Tyr
      65      70      75      80
Asn Arg Ser Ile Lys Asp Ile Lys Ala Ile Ser
      85      90

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<210> 223
 <211> 187
 <212> PRT
 <213> Pinus radiata

```

      <400> 223
Ser Phe Glu Ile His Thr Gly Lys Ser Ala Asp Ile Ser Arg Ala Gln
 1      5      10      15
Ser Ala Tyr Thr Gln Gln Asn Asn Asn Ile Phe Thr Ser Ser Lys Ile
      20      25      30
His Pro Val Val Ile Val Pro Gly Thr Gly Gly Asn Gln Val Glu Ala
      35      40      45
Arg Leu Thr Ala Asp Tyr Lys Pro Ser Gly Leu Leu Cys Arg Arg Trp
      50      55      60
Asn Trp Glu Arg Glu Trp Phe Arg Ile Trp Phe Asp Val Pro Val Val
      65      70      75      80
Leu Pro Pro Leu Thr Gln Cys Phe Ala Asp Arg Ile Ser Leu Val Tyr
      85      90      95
Asp Pro His Thr Asp Glu Tyr Tyr Asn Ala Pro Gly Val Glu Thr Arg
      100      105      110
Val Pro Tyr Phe Gly Ser Thr Glu Gly Met Lys Tyr Leu Asp Pro Cys
      115      120      125
Phe Lys Tyr Ile Thr Pro Tyr Met Ser Ser Leu Val Lys Ser Leu Glu
      130      135      140
Asp Val Gly Tyr Val Asp Gly Lys Ser Leu Phe Gly Ala Pro Tyr Asp
      145      150      155      160
Phe Arg Tyr Gly Pro Gly Thr Lys Ser Ser Ser Val Gly Ala Lys Tyr
      165      170      175

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Leu Glu Asn Leu Arg Lys Leu Val Glu Glu Ala
 180 185

<210> 224
 <211> 117
 <212> PRT
 <213> Pinus radiata

<400> 224
 Ser Ala Leu Ile Ile Gly Ser Phe Ile Phe Cys Ile Phe Leu Tyr Ile
 1 5 10 15
 Lys Gly His Val Ala Pro Ser Ser Thr Asp Ser Gly Ser Ser Gly Asn
 20 25 30
 Val Val Ile Asp Phe Tyr Trp Gly Met Glu Leu Tyr Pro Arg Ile Gly
 35 40 45
 Lys Asn Phe Asp Ile Lys Val Phe Thr Asn Cys Arg Phe Gly Met Met
 50 55 60
 Ser Trp Ala Val Leu Ala Val Thr Tyr Ser Ile Lys Gln Tyr Glu Glu
 65 70 75 80
 Tyr Gly Arg Val Ala Asp Ser Met Leu Val Ser Ser Ile Leu Met Val
 85 90 95
 Val Tyr Val Thr Lys Val Leu Leu Val Gly Ile Trp Leu Leu Glu His
 100 105 110
 His Gly Tyr Asn Ser
 115

<210> 225
 <211> 210
 <212> PRT
 <213> Pinus radiata

<400> 225
 Phe Ala Val Val Gly Pro Leu Gln Leu Thr Ser Tyr Pro Leu Ile Lys
 1 5 10 15
 Leu Val Gly Ile Arg Thr Gly Leu Pro Leu Pro Ser Leu Trp Glu Ile
 20 25 30
 Phe Ala Gln Leu Ala Val Tyr Phe Met Val Glu Asp Tyr Gly Asn Tyr
 35 40 45
 Trp Ile His Arg Trp Leu His Cys Lys Trp Gly Tyr Glu Lys Ile His
 50 55 60
 His Val His His Glu Phe Thr Ala Pro Met Gly Phe Ala Ala Pro Tyr
 65 70 75 80
 Ala His Trp Ser Glu Val Leu Ile Leu Gly Ile Pro Thr Phe Val Gly
 85 90 95
 Pro Ala Ile Ala Pro Gly His Met Ile Thr Phe Trp Cys Trp Val Val
 100 105 110
 Leu Arg Gln Val Glu Ala Ile Glu Thr His Ser Gly Tyr Asp Phe Pro
 115 120 125
 Trp Thr Leu Thr Lys Leu Ile Pro Phe Tyr Gly Gly Ala Glu Tyr His
 130 135 140
 Asp Tyr His His Tyr Val Gly Gly Gln Ser Gln Ser Asn Phe Ala Ser
 145 150 155 160
 Val Phe Thr Tyr Cys Asp Tyr Leu Tyr Gly Thr Asp Lys Gly Tyr Arg
 165 170 175
 Tyr Arg Lys Glu His Leu Leu Lys Ala Arg Glu Phe Glu Tyr Arg Leu
 180 185 190
 Lys Gln Met Ile Leu Arg Lys Lys Thr Ala Met Glu Gln Phe Gln Ile
 195 200 205
 Ser Leu
 210

<210> 226
 <211> 86
 <212> PRT
 <213> Pinus radiata

<400> 226
 Gly Pro His Leu Phe Thr Leu Trp Leu Trp Met Ser Leu Arg Val Leu
 1 5 10 15
 Glu Thr Val Glu Ala His Cys Gly Tyr Asp Phe Pro Trp Ser Ile Ser
 20 25 30
 Lys Leu Phe Pro Leu Tyr Gly Gly Ala Asp Phe His Asp Tyr His His
 35 40 45
 Arg Leu Leu Tyr Thr Lys Ser Gly Asn Tyr Ser Ser Thr Phe Thr Tyr
 50 55 60
 Met Asp Trp Leu Phe Gly Thr Asp Lys Gly Tyr Arg Lys Leu Lys Gly
 65 70 75 80
 Leu Gln Lys Asp Ser Lys
 85

<210> 227
 <211> 141
 <212> PRT
 <213> Pinus radiata

<400> 227
 Met Ala Thr Leu Val Glu Arg Gly Trp Leu Tyr Leu Ile Thr Asn Phe
 1 5 10 15
 Thr Asp Phe Gln Leu Ala Ser Ile Gly Ser Phe Leu Leu His Glu Ser
 20 25 30
 Ile Phe Tyr Leu Ser Gly Leu Pro Phe Ile Leu Leu Glu Thr Thr Gly
 35 40 45
 Leu Leu Ser Lys Tyr Lys Ile Gln Ser Lys Thr Asn Thr Val Ala Ala
 50 55 60
 Gln Glu Lys Cys Ile Thr Arg Leu Leu Leu Tyr His Phe Cys Val Asn
 65 70 75 80
 Leu Pro Val Met Val Val Ser Tyr Pro Val Phe Arg Phe Met Gly Met
 85 90 95
 Thr Ser Val Leu Pro Leu Pro Ser Trp Lys Val Val Val Ser Gln Leu
 100 105 110
 Val Cys Tyr Phe Ile Leu Glu Asp Phe Val Phe Tyr Trp Gly His Arg
 115 120 125
 Ile Leu His Ser Lys Trp Leu Tyr Lys His Val His Ser
 130 135 140

<210> 228
 <211> 381
 <212> PRT
 <213> Pinus radiata

<400> 228
 Met Gly Glu Glu Leu Gln Thr Trp Ile Leu Met Val Thr Ala Arg Ala
 1 5 10 15
 Pro Thr Asn Ile Ala Val Ile Lys Tyr Trp Gly Lys Arg Asp Glu Lys
 20 25 30
 Leu Ile Leu Pro Ile Asn Asp Ser Ile Ser Phe Thr Leu Asp Pro Asp
 35 40 45
 His Leu Ser Ala Thr Thr Thr Val Ala Val Ser Pro Ser Phe Thr Ser
 50 55 60
 Asp Arg Met Trp Leu Asn Gly Lys Glu Val Ser Leu Gly Gly Glu Arg
 65 70 75 80
 Tyr Gln Asn Cys Leu Arg Glu Ile Arg Ser Arg Gly Asn Asp Val Val

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<210> 229
<211> 81
<212> PRT
<213> Pinus radiata
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<210> 230
<211> 189
<212> PRT
<213> Pinus radiata
```

<400> 230

```

Met Pro Leu Thr Leu Leu Leu Ala Asn Thr Trp Ala Ser Ser Ala Ile
 1          5          10          15
Val Ser Arg Arg Val Ser Leu Phe Val Ala Cys Ser Thr Thr Val Val
          20          25          30
Ser Arg Ser Phe Ser Lys Ser Cys Ser Gly Ala Ile Pro Arg Lys Pro
          35          40          45
Lys Ser Ala His Pro Ala Leu Thr Gly Ser Arg Thr Cys Phe Ser Arg
          50          55          60
Asn Pro Ile Val Arg Asn Leu Ile Gly Ser Ala Ser Lys Met Gly Ala
65          70          75          80
Thr Val Glu Asp Thr Thr Met Asp Ala Val Gln Arg Arg Leu Met Phe
          85          90          95
Glu Asp Glu Cys Ile Leu Val Asp Glu Glu Asp His Val Ile Gly His
          100          105          110
Asp Ser Lys Tyr Asn Cys His Leu Met Glu Lys Ile Glu Ser Glu Asn
          115          120          125
Leu Leu His Arg Ala Phe Ser Val Phe Leu Phe Asn Thr Lys Tyr Glu
          130          135          140
Leu Leu Leu Gln Gln Arg Ser Ala Thr Lys Val Thr Phe Pro Leu Val
          145          150          155          160
Trp Thr Asn Thr Cys Cys Ser His Pro Leu Tyr Arg Glu Ser Glu Leu
          165          170          175
Ile Glu Glu Asn Asn Leu Gly Ser Glu Met Gln Pro Lys
          180          185

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<210> 231

<211> 113

<212> PRT

<213> Pinus radiata

<400> 231

```

Met Ala Gly Ile Pro Val Leu Arg Pro Phe Cys Ile Cys Leu Leu Ser
 1          5          10          15
Val Tyr Met Leu His Ile Val Ala Ala Val Ala Ser Pro Arg Leu Gly
          20          25          30
Arg Ser Ser Phe Pro Arg Gly Phe Lys Phe Gly Ala Gly Ser Ser Ala
          35          40          45
Tyr Gln Ala Glu Gly Ala Ala His Glu Gly Gly Lys Gly Pro Ser Ile
          50          55          60
Trp Asp Thr Phe Ser His Thr Pro Gly Lys Ile Ala Asp Gly Lys Asn
65          70          75          80
Gly Asp Val Ala Val Asp Gln Tyr His Arg Tyr Lys Glu Asp Val Gln
          85          90          95
Leu Leu Lys Tyr Met Gly Met Asp Val Tyr Arg Phe Ser Ile Ser Trp
          100          105          110
Ser

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<210> 232

<211> 127

<212> PRT

<213> Pinus radiata

<400> 232

```

Gly Pro Ser Ile Trp Asp Thr Phe Ser His Thr Pro Gly Lys Ile Ala
 1          5          10          15
Asp Gly Lys Asn Gly Asp Val Ala Val Asp Gln Tyr His Arg Tyr Lys
          20          25          30
Glu Asp Val Gln Leu Leu Lys Asn Met Gly Met Asp Val Tyr Arg Phe

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<210> 233
<211> 118
<212> PRT
<213> Eucalyptus grandis
```

```
<210> 234
<211> 111
<212> PRT
<213> Pinus radiata
```

```
<210> 235
<211> 391
<212> PRT
<213> Pinus radiata
```

<400> 235

Met Met Gln Lys Tyr Ile Gly Ala Asp Val Thr Ser Met Val Thr Leu
 1 5 10 15
 Pro Val Ile Ile Phe Glu Pro Met Thr Met Leu Gln Lys Ser Ala Glu
 20 25 30
 Leu Met Glu Tyr Thr Tyr Leu Leu Asp Met Ala Asp Glu Cys Glu Asp
 35 40 45
 Pro Tyr Leu Lys Met Ala Tyr Ala Ala Ser Trp Ala Ile Ser Val Tyr
 50 55 60
 Pro Ala Tyr Gln Arg Ser Trp Lys Pro Phe Asn Pro Ile Leu Gly Glu
 65 70 75 80
 Thr Tyr Glu Met Val Asn His Gly Gly Ile Thr Phe Ile Ala Glu Gln
 85 90 95
 Val Ser His His Pro Pro Met Gly Ser Ala Tyr Ala Glu Asn Glu His
 100 105 110
 Phe Thr Tyr Ser Leu Ser Ser Lys Val Lys Thr Lys Phe Leu Gly Asn
 115 120 125
 Ser Val Asp Ile Tyr Pro Leu Gly Arg Thr Arg Val Val Leu Lys Lys
 130 135 140
 Ser Gly Asp Val Leu Asp Leu Val Pro Pro Pro Ser Lys Val His Asn
 145 150 155 160
 Leu Ile Phe Gly Arg Thr Trp Ile Asp Ser Pro Gly Glu Met Val Leu
 165 170 175
 Thr Asn Leu Thr Thr Gly Asp Lys Val Val Leu Tyr Phe Gln Pro Cys
 180 185 190
 Gly Trp Phe Gly Ala Gly Arg Tyr Glu Val Asp Gly Tyr Val Tyr Asp
 195 200 205
 Ser Lys Glu Glu Pro Lys Ile Leu Met Thr Gly Lys Trp Asn Arg Ser
 210 215 220
 Met Gly Tyr Gln Pro Cys Asp Ala Glu Gly Glu Pro Leu Pro Gly Thr
 225 230 235 240
 Glu Leu Lys Glu Val Trp Arg Val Ala Asp Leu Pro Lys Asn Asp Lys
 245 250 255
 Phe Gln Tyr Thr Tyr Phe Ala His Lys Ile Asn Ser Phe Asp Thr Ala
 260 265 270
 Pro Lys Lys Leu Leu Ala Ser Asp Ser Arg Leu Arg Pro Asp Arg Ser
 275 280 285
 Ala Leu Glu Met Gly Asp Leu Ser Lys Ala Gly Ala Glu Lys Ser Asn
 290 295 300
 Leu Glu Glu Arg Gln Arg Ala Glu Lys Arg Cys Arg Glu Glu Lys Asn
 305 310 315 320
 Gln Pro Phe Thr Pro Arg Trp Phe Thr Val Thr Gly Glu Val Ala Thr
 325 330 335
 Thr Pro Trp Gly Asp Leu Glu Val Tyr Glu Tyr Asn Gly Lys Tyr Ser
 340 345 350
 Glu His Arg Ala Ser Val Asp Asp Ser Asn Phe Asp Asp Gly Thr Asp
 355 360 365
 Ser Lys Ser Met Glu Phe Asn Pro Trp Gln Tyr Gly Asn Ile Glu Ser
 370 375 380
 Gly Ser Asn Lys Lys Val Glu
 385 390

<210> 236

<211> 27

<212> PRT

<213> Pinus radiata

<400> 236

Met Met Gln Lys Tyr Ile Gly Ala Asp Val Thr Ser Met Val Thr Leu
 1 5 10 15
 Pro Val Ile Ile Phe Glu Pro Met Thr Met Leu

20

25

<210> 237
 <211> 134
 <212> PRT
 <213> Pinus radiata

<400> 237

Tyr	Leu	Val	Leu	Ile	Ser	Gln	Leu	Arg	Val	Gly	Met	Asp	Leu	Ser	Lys
1				5					10					15	
Val	Thr	Phe	Pro	Thr	Phe	Val	Leu	Glu	Pro	Arg	Ser	Met	Leu	Glu	Arg
			20					25					30		
Ile	Thr	Asp	Phe	Met	Ser	His	Pro	Asp	Leu	Ile	Phe	Gly	Ala	Glu	Asn
		35					40					45			
Ser	Asn	Asp	Pro	Glu	Glu	Arg	Phe	Met	Arg	Val	Leu	Ser	Tyr	Tyr	Leu
	50					55					60				
Ala	Gly	Trp	His	Ile	Lys	Pro	Lys	Gly	Val	Lys	Lys	Pro	Tyr	Asn	Pro
65					70				75					80	
Val	Leu	Gly	Glu	Phe	Phe	Arg	Cys	Arg	Tyr	Asp	Tyr	Ser	Asn	Asn	Thr
				85					90					95	
Gln	Gly	Phe	Tyr	Ile	Ala	Glu	Gln	Val	Ser	His	His	Pro	Pro	Ile	Ser
			100					105					110		
Ala	Phe	Phe	Tyr	Ile	Ser	Pro	Ala	Asn	Arg	Val	Ser	Ile	Ile	Gly	Glu
	115						120					125			
Leu	Arg	Pro	Lys	Ser	Lys										
130															

<210> 238
 <211> 133
 <212> PRT
 <213> Eucalyptus grandis

<400> 238

Ser	Ser	Lys	Gly	Arg	His	Cys	Lys	Pro	Phe	Asn	Pro	Leu	Leu	Gly	Glu
1				5					10					15	
Thr	Tyr	Glu	Ala	Asp	Tyr	Pro	Glu	Arg	Gly	Val	His	Phe	Phe	Ser	Glu
			20					25					30		
Lys	Val	Ser	His	His	Pro	Thr	Leu	Ile	Ala	Cys	His	Cys	Glu	Gly	Arg
		35					40					45			
Gly	Trp	Lys	Phe	Trp	Ala	Asp	Ser	Asn	Leu	Arg	Thr	Lys	Phe	Trp	Gly
	50					55					60				
Gln	Ser	Ile	Gln	Leu	Asp	Pro	Val	Gly	Ala	Leu	Thr	Leu	Glu	Phe	Asp
65					70				75					80	
Asp	Gly	Glu	Ile	Phe	Gln	Trp	Asn	Lys	Val	Thr	Thr	Ser	Ile	Asn	Asn
				85					90					95	
Leu	Ile	Ile	Gly	Lys	Val	Tyr	Cys	Asp	His	His	Gly	Val	Met	Asn	Ile
			100					105					110		
His	Gly	Asn	His	Gln	Tyr	Ser	Cys	Lys	Leu	Lys	Phe	Lys	Glu	Pro	Ser
	115						120					125			
Ile	Leu	Ala	Glu	Leu											
130															

<210> 239
 <211> 116
 <212> PRT
 <213> Eucalyptus grandis

<400> 239

Arg	Thr	Cys	Asp	Trp	Ser	Met	Arg	Ala	Ser	Trp	Ala	Ile	Ser	Val	Tyr
1				5					10					15	
Tyr	Ala	Tyr	Gln	Arg	Thr	Trp	Lys	Pro	Phe	Asn	Pro	Ile	Leu	Gly	Glu

20 25 30
 Thr Tyr Glu Leu Ala Asn His Gly Gly Ile Thr Phe Ile Ala Glu Gln
 35 40 45
 Val Cys His His Pro Pro Met Ser Ala Gly His Ala Glu Asn Asp His
 50 55 60
 Phe Thr Tyr Asp Val Thr Ser Lys Leu Lys Thr Lys Phe Leu Gly Asn
 65 70 75 80
 Ser Val Asp Val Tyr Pro Val Gly Arg Thr Arg Val Thr Leu Lys Arg
 85 90 95
 Asp Gly Val Val Leu Asp Leu Val Pro Pro Thr Lys Val Asn Asn
 100 105 110
 Leu Ile Phe Gly
 115

<210> 240
 <211> 105
 <212> PRT
 <213> Eucalyptus grandis

<400> 240
 Ser Arg Leu Arg Pro Asp Arg Tyr Ala Leu Glu Pro Gly Asp Leu Pro
 1 5 10 15
 Lys Ala Gly Ala Glu Lys Ser Ser Leu Glu Glu Arg Gln Arg Gly Glu
 20 25 30
 Lys Lys Asn Arg Glu Met Lys Gly Gln Lys Phe Thr Pro Arg Trp Phe
 35 40 45
 Asp Leu Thr Asp Glu Ile Ser Pro Thr Pro Trp Gly Asp Leu Glu Val
 50 55 60
 Tyr Arg Tyr Asn Gly Lys Tyr Thr Glu His Arg Ala Val Val Asp Ser
 65 70 75 80
 Leu Asp Thr Ile Glu Glu Ser Asp Ile Gln Ser Thr Glu Phe Asn Pro
 85 90 95
 Trp Gln Tyr Glu Ala Thr Phe Ala Glu
 100 105

<210> 241
 <211> 117
 <212> PRT
 <213> Pinus radiata

<400> 241
 Val Leu Arg Gly Leu Asp Thr Val Glu Asp Asp Thr Ser Ile Pro Leu
 1 5 10 15
 Asp Thr Lys Leu Pro Ile Leu Lys Ala Phe Tyr Lys His Ile Tyr Asp
 20 25 30
 Pro Ser Trp His Phe Ser Cys Gly Val Glu His Tyr Lys Glu Leu Met
 35 40 45
 Glu Lys Phe His His Val Ser Thr Thr Phe Leu Arg Leu Gly Arg Gly
 50 55 60
 Tyr Gln Glu Ala Ile Glu Glu Ile Thr Lys Lys Met Gly Ala Gly Met
 65 70 75 80
 Ala Lys Phe Ile Cys Lys Glu Val Glu Ser Val Glu Asp Tyr Asp Glu
 85 90 95
 Tyr Cys His Tyr Val Ala Gly Leu Val Gly Phe Gly Leu Ser Arg Leu
 100 105 110
 Phe His Ala Ala Gln
 115

<210> 242
 <211> 190
 <212> PRT

<213> Pinus radiata

<400> 242

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Met Ala Ile Tyr Thr Pro Gln Pro Ala His Arg Leu Ile Ser Trp Ser
 1          5          10          15
Thr Met Glu Asn His Thr Val Val Ile Ala Ala Ala Ile Ser Phe Val
 20          25          30
Ser Val Leu Leu Ser Tyr Tyr Ile Val Leu Ser Arg Trp Lys Arg Arg
 35          40          45
Ser Asn Gly Leu Arg Gly Ile Gln Ser Lys Ser Phe Glu Lys Ser Thr
 50          55          60
Asp Asp Asn Gly Ile Ala Ile Glu Ala Ala Gly Gly Thr Asp Val Ile
 65          70          75          80
Ile Val Gly Ala Gly Val Ala Gly Ser Ala Leu Ala Tyr Thr Leu Gly
 85          90          95
Lys Asp Gly Arg Arg Ile His Val Ile Glu Arg Asp Leu Ser Glu Pro
 100          105          110
Asp Arg Ile Val Gly Glu Leu Leu Gln Pro Gly Gly Tyr Leu Lys Leu
 115          120          125
Ile Glu Leu Gly Leu Gln Asp Cys Val Glu Gly Ile Asp Ala Gln Ser
 130          135          140
Ile Phe Gly Asp Ala Leu Phe Lys Glu Gly Lys Asp Thr Lys Val Ala
 145          150          155          160
Tyr Pro Leu Glu Asn His His Ala Asp Arg Ala Gly Arg Ser Phe His
 165          170          175
Asn Gly Arg Phe Ile Gln Arg Met Arg Glu Lys Ala Ala Ser
 180          185          190

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<210> 243

<211> 124

<212> PRT

<213> Pinus radiata

<400> 243

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Cys Leu Thr Thr Asp Ser Gly Gln Val Ile Asn Cys Arg Asn Arg Tyr
 1          5          10          15
Thr Ala Met Ala Ile Tyr Thr Pro Gln Pro Ala His Arg Leu Ile Ser
 20          25          30
Trp Ser Thr Met Glu Asn His Thr Val Ala Ile Ala Val Ala Ile Gly
 35          40          45
Phe Val Ser Val Leu Leu Ser Tyr Tyr Ile Val Leu Asn Arg Trp Lys
 50          55          60
Arg Arg Ser Asn Gly Leu Arg Gly Ile Gln Ser Lys Ser Phe Glu Lys
 65          70          75          80
Ser Thr Asp Asp Asn Gly Ile Ala Ile Glu Ala Ala Gly Gly Thr Asp
 85          90          95
Val Ile Ile Val Gly Ala Gly Val Ala Gly Ser Ala Leu Ala Tyr Thr
 100          105          110
Leu Gly Lys Asp Gly Arg Arg Ile His Val Ile Glu
 115          120

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<210> 244

<211> 123

<212> PRT

<213> Eucalyptus grandis

<400> 244

```

Met Asp Gly Gln Tyr Leu Val Ser Gly Val Leu Ala Leu Phe Leu Gly
 1          5          10          15
Ile Phe Leu Leu Tyr Lys Gly Leu Gly Lys Gln Lys Arg Arg Leu Ser
 20          25          30

```

Lys Lys Gly Arg Gly Asp Asp Tyr Val Lys Ser Ser Val Asp Gly Gly
 35 40 45
 Phe Val Pro Gly Gly Val Asp Gly Ser Thr Asp Ile Val Ile Val Gly
 50 55 60
 Ala Gly Val Ala Gly Ala Ala Leu Ala Tyr Ala Leu Gly Lys Asp Gly
 65 70 75 80
 Arg Arg Val Arg Val Ile Glu Arg Asp Leu Thr Glu Gln Asp Arg Ile
 85 90 95
 Val Gly Glu Leu Leu Gln Pro Gly Gly Tyr Leu Lys Leu Met Glu Leu
 100 105 110
 Asp Leu Ala Asp Cys Val Gln Thr Ile Asp Ala
 115 120

<210> 245
 <211> 221
 <212> PRT
 <213> Eucalyptus grandis

<400> 245
 Leu Gly Ser Lys Tyr Lys Pro Gln Glu Glu Phe Val Glu Trp Ile Gln
 1 5 10 15
 Lys Gly Thr Lys Pro Ile Tyr Ile Gly Phe Gly Ser Met Pro Leu Glu
 20 25 30
 Asp Pro Lys Lys Thr Thr Asp Ile Ile Ile Lys Ala Leu Thr Asp Thr
 35 40 45
 Gly Gln Arg Gly Ile Val Gly Arg Gly Trp Gly Asp Leu Gly Thr Leu
 50 55 60
 Leu Asp Val Pro Asp Ser Val Phe Leu Leu Glu Asp Cys Pro His Asp
 65 70 75 80
 Trp Leu Phe Pro Gln Cys Ser Ala Val Val His His Gly Gly Ala Gly
 85 90 95
 Thr Thr Ala Thr Gly Leu Lys Ala Gly Cys Pro Thr Thr Ile Val Pro
 100 105 110
 Phe Phe Gly Asp Gln Phe Phe Trp Gly Asp Arg Val His Gln Arg Gly
 115 120 125
 Leu Gly Pro Ala Pro Ile Pro Ile Ser Gln Leu Ser Val Glu Asn Leu
 130 135 140
 Ser Asp Ala Ile Arg Phe Met Leu Gln Pro Glu Val Lys Ser Gln Ala
 145 150 155 160
 Met Glu Met Ala Lys Leu Ile Glu Asn Glu Asp Gly Val Ala Ala Ala
 165 170 175
 Val Asp Ala Phe His Arg His Leu Pro Glu Glu Phe Pro Ser Ser Ser
 180 185 190
 Val Ser Ser Asp Gly Glu Glu His Pro Asn Pro Phe Leu Trp Leu Phe
 195 200 205
 Leu Gln Val Glu Lys Trp Cys Cys Leu Pro Cys Ser Lys
 210 215 220

<210> 246
 <211> 114
 <212> PRT
 <213> Eucalyptus grandis

<400> 246
 Leu Asp Asn Cys Pro His Asp Trp Leu Phe Leu Arg Cys Ser Ala Val
 1 5 10 15
 Val His His Gly Gly Ala Gly Thr Thr Ala Ala Gly Leu Lys Ala Ala
 20 25 30
 Cys Pro Thr Thr Val Val Pro Phe Phe Gly Asp Gln Pro Phe Trp Gly
 35 40 45
 Glu Arg Val His Ala Arg Gly Val Gly Pro Val Pro Ile Pro Val Asp

50 55 60
 Glu Phe Ser Leu Glu Lys Leu Val Asp Ala Ile Arg Phe Met Leu Asp
 65 70 75 80
 Pro Lys Val Lys Gln Cys Ala Glu Glu Leu Ala Lys Asp Met Glu His
 85 90 95
 Glu Asp Gly Val Glu Gly Ala Val Lys Ala Phe Tyr Lys His Phe Pro
 100 105 110
 Arg Glu

<210> 247
 <211> 140
 <212> PRT
 <213> Pinus radiata

<400> 247
 Met Ala Thr Gly Gly Gly Ala Leu Asp Leu Ala Ser Gly Met Gly Gly
 1 5 10 15
 Asn Ile Glu Lys Glu Gln Met Leu Thr Ala Val Glu Glu Tyr Glu Lys
 20 25 30
 Tyr His Met Tyr Tyr Gly Gly Asp Glu Gly Ser Arg Lys Ser Asn Tyr
 35 40 45
 Thr Asp Met Val Asn Lys Tyr Tyr Asp Leu Ala Thr Ser Phe Tyr Glu
 50 55 60
 Tyr Gly Trp Gly Glu Ser Phe His Phe Ala His Arg Trp Lys Gly Glu
 65 70 75 80
 Thr Leu Arg Glu Ser Ile Lys Arg His Glu His Phe Leu Ala Leu His
 85 90 95
 Leu Cys Leu Lys Pro Ala Met Lys Val Leu Asp Val Gly Cys Gly Ile
 100 105 110
 Gly Gly Pro Leu Arg Glu Ile Ala Arg Phe Ser Arg Thr Ser Ile Thr
 115 120 125
 Gly Leu Asn Asn Asn Ala Tyr Gln Ile Ser Arg Gly
 130 135 140

<210> 248
 <211> 152
 <212> PRT
 <213> Eucalyptus grandis

<400> 248
 Met Ser Lys Ala Gly Ala Met Asp Leu Ala Thr Gly Leu Gly Gly Lys
 1 5 10 15
 Met Asp Lys Ser Asp Val Leu Ser Ala Val Asp Lys Tyr Glu Lys Tyr
 20 25 30
 His Val Cys Tyr Gly Gly Asp Glu Glu Glu Arg Arg Ala Asn Tyr Ser
 35 40 45
 Asp Met Val Asn Lys Tyr Tyr Asp Leu Ala Thr Ser Phe Tyr Glu Phe
 50 55 60
 Gly Trp Gly Glu Ser Phe His Phe Ala His Arg Trp Lys Gly Glu Ser
 65 70 75 80
 Leu Arg Glu Ser Ile Lys Arg His Glu His Phe Leu Ala Leu Gln Leu
 85 90 95
 Gly Leu Lys Pro Gly His Lys Val Leu Asp Val Gly Cys Gly Ile Gly
 100 105 110
 Gly Pro Leu Arg Glu Ile Ala Arg Phe Ser Ser Ala Ser Val Thr Gly
 115 120 125
 Leu Asn Asn Asn Glu Tyr Gln Ile Thr Arg Gly Lys Glu Leu Asn Arg
 130 135 140
 Ile Ala Gly Val Asp Lys Thr Cys
 145 150

<210> 249
 <211> 100
 <212> PRT
 <213> Eucalyptus grandis

<400> 249
 Met Ser Lys Ala Gly Ala Met Asp Leu Ala Thr Gly Leu Gly Gly Lys
 1 5 10 15
 Met Asp Lys Ser Asp Val Leu Ser Ala Val Asp Lys Tyr Glu Lys Tyr
 20 25 30
 His Val Cys Tyr Gly Gly Asp Glu Glu Glu Arg Arg Ala Asn Tyr Ser
 35 40 45
 Asp Met Val Asn Lys Tyr Tyr Asp Leu Ala Thr Ser Phe Tyr Glu Phe
 50 55 60
 Gly Trp Gly Glu Ser Phe His Phe Ala His Arg Trp Lys Gly Glu Ser
 65 70 75 80
 Leu Arg Glu Ser Ile Lys Arg His Glu His Phe Leu Ala Leu Gln Leu
 85 90 95
 Gly Leu Lys Pro
 100

<210> 250
 <211> 148
 <212> PRT
 <213> Eucalyptus grandis

<400> 250
 Ala Met Pro Trp Tyr Cys Ala Leu Pro Thr Leu Ser Glu Tyr Met Val
 1 5 10 15
 Glu Asn Gly Trp Thr Lys Cys Phe Ser Arg Ile Ser Asp Val Gly Trp
 20 25 30
 Leu Ala Tyr Leu Val Tyr Leu Ser Ile Tyr Leu Val Met Ala Glu Phe
 35 40 45
 Gly Ile Tyr Trp Met His Arg Glu Leu His Asp Ile Lys Pro Leu Tyr
 50 55 60
 Lys His Leu His Ala Thr His His Ile Tyr Asn Lys Gln Asn Thr Leu
 65 70 75 80
 Ser Pro Phe Ala Gly Leu Ala Phe His Pro Leu Asp Gly Ile Leu Gln
 85 90 95
 Ala Val Pro His Val Met Ala Leu Phe Leu Val Pro Thr His Phe Thr
 100 105 110
 Thr His Ile Ala Leu Leu Phe Leu Glu Ala Ile Trp Thr Ala Asn Ile
 115 120 125
 His Asp Cys Ile His Gly Lys Leu Trp Pro Val Met Gly Ala Gly Tyr
 130 135 140
 His Thr Ile His
 145

<210> 251
 <211> 201
 <212> PRT
 <213> Eucalyptus grandis

<400> 251
 Phe Met Ser Cys Leu Pro Asn Met Ile Val Met Ala Pro Ser Asp Glu
 1 5 10 15
 Asp Glu Leu Val Asp Met Val Glu Thr Ala Ala Ile Val Asp Asp Arg
 20 25 30
 Pro Ile Cys Phe Arg Tyr Pro Arg Gly Ala Ile Val Arg Thr Asp Lys
 35 40 45

Ser Leu Ser Gln Gly Ile Pro Ile Glu Ile Gly Lys Gly Arg Ile Leu
 50 55 60
 Ala Glu Gly Lys Asp Val Ala Leu Leu Gly Tyr Gly Ser Met Val Gln
 65 70 75 80
 Asn Cys Val Lys Ala Arg Ser Leu Leu Ser Lys Leu Gly Ile Glu Val
 85 90 95
 Thr Val Ala Asp Ala Arg Phe Cys Lys Pro Leu Asp Ile Gly Leu Leu
 100 105 110
 Arg Glu Leu Cys Glu Asn His Ala Phe Leu Val Thr Val Glu Glu Gly
 115 120 125
 Ser Ile Gly Gly Phe Gly Ser His Val Ala Gln Phe Ile Ala Leu Asp
 130 135 140
 Gly Arg Leu Asp Gly Arg Ile Lys Trp Arg Pro Ile Val Leu Pro Asp
 145 150 155 160
 Ala Tyr Val Glu His Ala Ser Pro Asn Glu Gln Leu Ser Leu Ala Gly
 165 170 175
 Leu Thr Gly His His Ile Ala Ala Thr Val Leu Ser Leu Leu Gly Arg
 180 185 190
 Thr Arg Glu Ala Leu Leu Leu Met Cys
 195 200

<210> 252
 <211> 138
 <212> PRT
 <213> Eucalyptus grandis

<400> 252
 Asp Ile Lys Lys Ile Val Glu Leu Met Ser Asp Leu His Phe Ile Tyr
 1 5 10 15
 Asn Thr His Arg Phe Ala Tyr Leu Tyr Ser Lys Phe Asn Ser Ser Ile
 20 25 30
 Tyr Met Tyr Lys Phe Ser Leu Asp Thr Asp Leu Asn Ile Val Lys Lys
 35 40 45
 Met Ser Gly Phe Asp Val Glu Gly Val Cys His Ala Asp Glu Leu Phe
 50 55 60
 Tyr Phe Phe Ser Thr Asn Met Thr Lys Asp Tyr Tyr Glu Ser Glu Asp
 65 70 75 80
 Lys Ile Lys Glu Tyr Val Trp Lys Val Thr Lys Leu Trp Thr Asn Phe
 85 90 95
 Ala Lys Thr Ser Asn Pro Thr Pro Asp Thr Ser Leu Gly Val Ser Trp
 100 105 110
 Pro Arg Tyr Thr Met Ala Asn Lys Glu Tyr Leu Asp Ile Asn Thr Gln
 115 120 125
 Leu Thr Thr Gly Arg Tyr Ser Glu Arg Glu
 130 135

<210> 253
 <211> 610
 <212> PRT
 <213> Pinus radiata

<400> 253
 Cys Leu Leu Leu Leu Gln Leu Lys Leu Phe Cys Ser Pro Ile Asn Met
 1 5 10 15
 Ala Ile Ala Ser Arg Ala Gly Val Ala Pro Ile Leu Gln Val Asp Cys
 20 25 30
 His Phe Thr His Phe Asn Ser Met Thr Glu Leu Gly Ser Arg Asn Ser
 35 40 45
 Met Met Phe Gln Ser Ala Ile Pro Cys Ser Phe Arg Gln Ile Arg Ala
 50 55 60
 Thr Thr Lys Arg Lys Arg Cys Val Leu Leu Ala Lys Leu Ser Asn Ser

65	70	75	80
Asp Gly Glu Asn Gly Lys Asn Val Lys Ala Ala Val Glu Ile Ala Ser			
	85	90	95
Lys Ser Gly Phe Pro Ala Glu Lys Pro Pro Thr Pro Leu Leu Asp Thr			
	100	105	110
Val Asn Tyr Pro Val His Leu Lys Asn Leu Ser Ile Gln Asp Leu Glu			
	115	120	125
Gln Leu Ala Thr Glu Ile Arg Ala Glu Leu Val Phe Gly Val Ala Lys			
	130	135	140
Thr Gly Gly His Leu Gly Gly Ser Leu Gly Val Val Asp Leu Thr Val			
	145	150	155
Ala Leu His His Val Phe Asp Ser Pro Glu Asp Arg Ile Ile Trp Asp			
	165	170	175
Val Gly His Gln Ser Tyr Pro His Lys Ile Leu Thr Gly Arg Arg Ser			
	180	185	190
Lys Met His Thr Ile Arg Gln Thr Ser Gly Leu Ala Gly Phe Pro Lys			
	195	200	205
Arg Asp Glu Ser Lys Tyr Asp Ala Phe Gly Ala Gly His Ser Ser Thr			
	210	215	220
Ser Ile Ser Ala Gly Leu Gly Met Ala Val Gly Arg Asp Leu Leu Lys			
	225	230	235
Lys Asn Asn His Val Val Ala Val Ile Gly Asp Gly Ala Met Thr Ala			
	245	250	255
Gly Gln Ala Tyr Glu Ala Met Asn Asn Ser Gly Tyr Leu Glu Ser Asn			
	260	265	270
Leu Ile Ile Ile Leu Asn Asp Asn Lys Gln Val Ser Leu Pro Thr Ala			
	275	280	285
Thr Leu Asp Gly Ala Ala Pro Val Gly Ala Leu Thr Arg Ala Leu			
	290	295	300
Thr Lys Leu Gln Ser Ser Lys Lys Leu Arg Lys Leu Arg Glu Ala Ala			
	305	310	315
Lys Gly Leu Thr Lys Gln Ile Gly Gly Pro Thr His Glu Val Ala Ser			
	325	330	335
Lys Val Asp Lys Tyr Ala Arg Gly Leu Ile Ser Pro Ala Ser Ser Ser			
	340	345	350
Leu Phe Asp Glu Leu Gly Leu Tyr Tyr Ile Gly Pro Val Asp Gly His			
	355	360	365
Asn Ile Glu Asp Met Val Thr Ile Leu Glu Lys Ile Lys Ser Met Pro			
	370	375	380
Ala Thr Gly Pro Val Leu Ile His Leu Val Thr Glu Lys Gly Lys Gly			
	385	390	395
Tyr Pro Pro Ala Glu Glu Ala Ala Asp Lys Leu His Gly Val Val Lys			
	405	410	415
Phe Asp Pro Val Thr Gly Lys Gln Phe Lys Ser Lys Ser Ser Val Leu			
	420	425	430
Ser Tyr Thr Gln Tyr Phe Ala Glu Ala Leu Ile Ala Glu Ala Glu Val			
	435	440	445
Asp Ser Lys Ile Val Ala Ile His Ala Ala Met Gly Gly Gly Thr Gly			
	450	455	460
Leu Asn Tyr Phe Gln Lys Lys Phe Pro Glu Arg Cys Phe Asp Val Gly			
	465	470	475
Ile Ala Glu Gln His Ala Val Thr Phe Ala Ala Gly Leu Ala Thr Glu			
	485	490	495
Gly Leu Lys Pro Phe Cys Ala Ile Tyr Ser Thr Phe Leu Gln Arg Gly			
	500	505	510
Tyr Asp Gln Val Val His Asp Val Asp Leu Gln Lys Leu Pro Val Arg			
	515	520	525
Phe Ala Met Asp Arg Ala Gly Leu Val Gly Ala Asp Gly Pro Thr His			
	530	535	540
Cys Gly Ser Phe Asp Val Ala Tyr Met Ala Cys Leu Pro Asn Met Ile			
545	550	555	560

Val Met Ala Pro Ser Asp Glu Val Glu Leu Met His Ile Val Ala Thr
 565 570 575
 Ala Ala Ala Ile Asp Asp Arg Pro Ser Cys Phe Arg Phe Pro Arg Gly
 580 585 590
 Asn Gly Val Gly Leu Ser Asn Leu Pro Leu Asn Asn Lys Gly Val Pro
 595 600 605
 Leu Glu
 610

<210> 254
 <211> 147
 <212> PRT
 <213> Eucalyptus grandis

<400> 254
 Met Ala Asp Leu Lys Ser Lys Phe Met Glu Ala Tyr Ala Val Leu Lys
 1 5 10 15
 Lys Glu Leu Leu Ala Asp Pro Ala Phe Glu Phe Ser Asp Glu Ser Arg
 20 25 30
 Gln Trp Val Asp Arg Met Leu Asp Tyr Asn Val Pro Gly Gly Lys Leu
 35 40 45
 Asn Arg Gly Leu Ser Val Ile Asp Ser Tyr Lys Leu Leu Lys Glu Gly
 50 55 60
 Lys Glu Leu Thr Glu Glu Glu Ile Phe Leu Ala Ser Ala Leu Gly Trp
 65 70 75 80
 Cys Ile Glu Trp Leu Gln Ala Tyr Phe Leu Val Leu Asp Asp Ile Met
 85 90 95
 Asp Ser Ser His Thr Arg Arg Gly Gln Pro Cys Trp Phe Arg Leu Pro
 100 105 110
 Lys Val Gly Met Ile Ala Ala Asn Asp Gly Val Leu Leu Arg Asn His
 115 120 125
 Ile Pro Arg Ile Leu Lys Asn His Phe Arg Gly Lys Pro Tyr Tyr Val
 130 135 140
 Asp Leu Leu
 145

<210> 255
 <211> 123
 <212> PRT
 <213> Eucalyptus grandis

<400> 255
 Phe Pro Leu Ser Ser Ser Ser Leu Cys Ser Glu Phe Pro Phe Cys Val
 1 5 10 15
 Ala Gly Arg Ala Arg Gln Ala Gly Ala Gly Trp Ala Gly Glu Ser
 20 25 30
 Ser Val Val Ala Ser Met Ala Asp Leu Asn Ser Lys Leu Leu Glu Ala
 35 40 45
 Asn Ala Val Leu Lys Lys Glu Leu Pro Glu Asp Pro Ala Phe Glu Phe
 50 55 60
 Ser Asp Asp Ser Arg Gln Trp Val Glu Arg Glu Asn Tyr Gly Lys Pro
 65 70 75 80
 Asp Ser Ala Asn Val Ala Lys Val Lys Val Leu Tyr His Glu Ile Asn
 85 90 95
 Leu Gln Gly Tyr Cys Lys Ser Ile Ser Lys Asn Lys Asn Ile Pro Thr
 100 105 110
 Val Lys Ala Asn Ala Asn Ser Val Glu Ala Thr
 115 120

<210> 256
 <211> 127

<212> PRT

<213> Pinus radiata

<400> 256

```

Arg Pro Cys His Leu Glu Trp Ile His Ile His Lys Thr Ala Val Ile
 1           5           10           15
Leu Glu Cys Ser Val Val Cys Gly Asp Ile Ile Ser Gly Ala Ser Glu
      20           25           30
Asn Glu Ile Glu Arg Ile Lys Ser Tyr Ala Arg Ser Val Gly Leu Leu
      35           40           45
Phe Gln Val Val Asp Asp Ile Leu Asp Val Thr Lys Ser Ser Lys Glu
 50           55           60
Leu Gly Lys Thr Ala Gly Lys Asp Leu Ile Thr Asp Lys Ala Thr Tyr
65           70           75           80
Pro Lys Leu Met Gly Leu Glu Thr Ala Lys Gln Phe Ala Val Glu Leu
      85           90           95
Leu Gly Arg Ala Lys Glu Asp Leu Ser Cys Phe Asp Pro Lys Lys Ala
      100           105           110
Ala Pro Leu Leu Gly Ile Ala Glu Tyr Ile Ala Phe Arg Gln Asn
      115           120           125

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<210> 257

<211> 196

<212> PRT

<213> Eucalyptus grandis

<400> 257

```

Ala Cys Ala Val Glu Met Ile His Thr Met Ser Leu Ile His Asp Asp
 1           5           10           15
Leu Pro Cys Met Asp Asn Asp Asp Leu Arg Arg Gly Lys Pro Thr Asn
      20           25           30
His Lys Val Tyr Gly Glu Asp Val Ala Val Leu Ala Gly Asp Ala Leu
      35           40           45
Leu Ala Tyr Ala Phe Glu His Ile Ala Val Glu Thr Lys Gly Val Ser
 50           55           60
Pro Thr Arg Ile Val Arg Ala Ile Phe Glu Leu Ala Arg Ser Ile Gly
65           70           75           80
Ala Glu Gly Leu Val Ala Gly Gln Val Val Asp Ile Ser Ser Glu Gly
      85           90           95
Ile Ala Asn Val Gly Leu Glu His Leu Glu Phe Ile His Leu His Lys
      100           105           110
Thr Ala Ala Leu Leu Glu Ala Ser Val Val Leu Gly Ala Ile Met Gly
      115           120           125
Gly Gly Ser Asn Glu Glu Val Lys Leu Arg Gly Phe Ala Arg Cys
      130           135           140
Ile Gly Leu Leu Phe Gln Val Val Asp Asp Ile Leu Asp Leu Thr Gln
145           150           155           160
Ser Ser Gln Glu Leu Gly Lys Thr Ala Gly Lys Asp Leu Val Ala Asp
      165           170           175
Lys Val Thr Tyr Pro Lys Leu Met Gly Ile Glu Lys Ser Arg Glu Leu
      180           185           190
Ala Asn Lys Leu
      195

```

<210> 258

<211> 159

<212> PRT

<213> Eucalyptus grandis

<400> 258

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Met Gly Ser Leu Gly Ala Ile Leu Lys His Pro Asp Asp Phe Tyr Pro

```

1	5	10	15
Leu Leu Lys	Leu Lys Ile Ala Ala	Arg Asn Ala Glu Lys Arg Ile Pro	
	20	25	30
Pro Gln Pro	His Trp Gly Phe Cys Tyr Ser Met Leu His Lys Val Ser		
	35	40	45
Arg Ser Phe	Gly Leu Val Ile Gln Gln Leu Gly Pro Glu Leu Arg Asp		
	50	55	60
Ala Val Cys	Ile Phe Tyr Leu Val Leu Arg Ala Leu Asp Thr Val Glu		
65	70	75	80
Asp Asp Thr	Ser Ile Pro Thr Asp Val Lys Val Pro Ile Leu Lys Ala		
	85	90	95
Phe His Gln	His Val Tyr Asp Lys Glu Trp His Phe Ser Cys Gly Thr		
	100	105	110
Lys Glu Tyr	Lys Val Leu Met Asp Gln Phe His His Val Ser Thr Ala		
	115	120	125
Phe Leu Glu	Leu Gly Lys Ser Tyr Gln Glu Ala Ile Asp Asp Ile Thr		
	130	135	140
Lys Arg Met	Gly Ala Gly Met Ala Lys Phe Ile Cys Gln Glu Val		
145	150	155	

<210> 259

<211> 106

<212> PRT

<213> Pinus radiata

<400> 259

Met Ala Ile Tyr Thr Pro Gln Pro Ala His Arg Leu Ile Ser Trp Ser	
1	5
Thr Met Glu Asn His Thr Val Ala Ile Ala Val Ala Ile Gly Phe Val	
	20
Ser Val Leu Leu Ser Tyr Tyr Ile Val Leu Asn Arg Trp Lys Arg Arg	
	35
Ser Asn Gly Leu Arg Gly Ile Gln Ser Lys Ser Phe Glu Lys Ser Thr	
	50
Asp Asp Asn Gly Ile Ala Ile Glu Ala Ala Gly Gly Thr Asp Val Ile	
65	70
Ile Val Gly Ala Gly Val Ala Gly Ser Ala Leu Ala Tyr Thr Leu Gly	
	85
Lys Asp Gly Arg Arg Ile His Val Ile Glu	
	100
	105

<210> 260

<211> 93

<212> PRT

<213> Pinus radiata

<400> 260

Met Ala Ile Tyr Thr Pro Gln Pro Ala His Arg Leu Ile Ser Trp Ser	
1	5
Thr Met Glu Asn His Thr Val Val Ile Ala Ala Ala Ile Ser Phe Val	
	20
Ser Val Leu Leu Ser Tyr Tyr Ile Val Leu Ser Arg Trp Lys Arg Arg	
	35
Ser Asn Gly Leu Arg Gly Ile Gln Ser Lys Ser Phe Glu Lys Ser Thr	
	50
Asp Asp Asn Gly Ile Ala Ile Glu Ala Ala Gly Gly Thr Asp Val Ile	
65	70
Ile Val Gly Ala Gly Val Ala Gly Ser Ala Leu Ala Tyr	
	85
	90

<210> 261

<211> 217
 <212> PRT
 <213> Eucalyptus grandis

<400> 261
 Pro Gln Leu Tyr Lys Ala Phe Ile Ala Ala Ile Asp Lys Gly Asn Ile
 1 5 10 15
 Lys Ser Met Pro Asn Arg Ser Met Pro Ala Asn Pro Gln Pro Thr Pro
 20 25 30
 Gly Ala Leu Leu Met Gly Asp Ala Phe Asn Met Arg His Pro Leu Thr
 35 40 45
 Gly Gly Gly Met Thr Val Ala Leu Ser Asp Ile Val Leu Leu Arg Asn
 50 55 60
 Leu Leu Arg Pro Leu Gln Asp Leu Asn Asp Ala Ser Ala Leu Cys Lys
 65 70 75 80
 Tyr Leu Glu Ser Phe Tyr Thr Leu Arg Lys Pro Val Ala Ser Thr Ile
 85 90 95
 Asn Thr Leu Ala Gly Ala Leu Tyr Lys Val Phe Cys Ala Ser Pro Asp
 100 105 110
 Pro Ala Arg Lys Glu Met Arg Gln Ala Cys Phe Asp Tyr Leu Ser Leu
 115 120 125
 Gly Gly Leu Cys Ser Thr Gly Pro Val Ser Leu Leu Ser Gly Leu Asn
 130 135 140
 Pro Arg Pro Met His Leu Val Cys His Phe Phe Ala Val Ala Val Tyr
 145 150 155 160
 Gly Val Gly Arg Leu Cys Leu Pro Phe Pro Ser Pro Lys Arg Met Trp
 165 170 175
 Leu Gly Ala Arg Leu Val Lys Gly Ala Ser Gly Ile Ile Phe Pro Ile
 180 185 190
 Ile Arg Asp Glu Gly Val Arg Gln Met Phe Phe Pro Ala Thr Val Pro
 195 200 205
 Ala Tyr His Arg Ala Pro Pro Val His
 210 215

<210> 262
 <211> 94
 <212> PRT
 <213> Eucalyptus grandis

<400> 262
 Met Glu Asp Asp Arg Asp Arg Gly Leu Leu Tyr Asp Ser Asp Pro Ser
 1 5 10 15
 Ser Ser Ser Leu Ser Pro Pro Arg Pro Phe Ala Leu Thr Phe Phe Asp
 20 25 30
 Arg Glu Arg His Val Thr Phe Leu Glu Met Met Tyr His Met Leu Pro
 35 40 45
 Arg Pro Tyr Gln Ser Gln Glu Ile Asn His Leu Thr Leu Ala Tyr Phe
 50 55 60
 Val Ile Ser Gly Leu Asp Ile Leu Asp Ala Leu Asp Arg Val His Lys
 65 70 75 80
 Asp Ala Val Ala Asp Trp Val Leu Ser Phe Gln Ala His Phe
 85 90

<210> 263
 <211> 81
 <212> PRT
 <213> Eucalyptus grandis

<400> 263
 Glu Ile Leu Thr Lys Val Ile Ser Leu Ala Ser Ile Met Asp Asp Ile
 1 5 10 15

Tyr Asp Val Tyr Gly Thr Leu Glu Glu Leu Ala Leu Leu Asn Glu Ala
 20 25 30
 Ile Gln Lys Trp Asp Phe Asp Ala Met Asp Gly Leu Pro Glu Tyr Met
 35 40 45
 Gln Ala Tyr Phe Lys Glu Phe Leu Gln Leu Tyr Glu Tyr Ile Gly Asn
 50 55 60
 Gln Leu Ala Ala Lys Gly Arg Ser Tyr Arg Leu Ile Tyr Ala Lys Glu
 65 70 75 80
 Val

<210> 264
 <211> 125
 <212> PRT
 <213> Pinus radiata

<400> 264
 Leu Tyr Arg Ala Ser Leu Ile Ala Phe Pro Gly Glu Lys Val Met Asp
 1 5 10 15
 Glu Ala Glu Thr Phe Ser Ala Lys Tyr Leu Lys Glu Ala Leu Gln Lys
 20 25 30
 Ile Pro Val Ser Ser Leu Ser Arg Glu Ile Gly Asp Val Leu Glu Tyr
 35 40 45
 Gly Trp His Thr Tyr Leu Pro Arg Leu Glu Ala Arg Asn Tyr Ile Asp
 50 55 60
 Val Phe Gly Gln Asp Thr Glu Asn Ser Lys Ser Tyr Met Lys Thr Glu
 65 70 75 80
 Lys Leu Leu Glu Leu Ala Lys Leu Glu Phe Asn Ile Phe His Ala Leu
 85 90 95
 Gln Lys Arg Glu Leu Glu Tyr Leu Val Arg Trp Trp Lys Gly Ser Gly
 100 105 110
 Ser Pro Gln Met Thr Phe Cys Arg His Arg His Val Glu
 115 120 125

<210> 265
 <211> 219
 <212> PRT
 <213> Pinus radiata

<400> 265
 Met Pro Gln Asp Met Lys Ile Cys Phe Lys Gly Phe Tyr Asn Thr Phe
 1 5 10 15
 Asn Glu Ile Ala Glu Glu Gly Arg Lys Arg Gln Gly Arg Asp Val Leu
 20 25 30
 Ser Tyr Ile Gln Lys Val Trp Glu Val Gln Leu Glu Ala Tyr Thr Lys
 35 40 45
 Glu Ala Glu Trp Ser Ala Val Arg Tyr Val Pro Ser Tyr Asp Glu Tyr
 50 55 60
 Ile Gly Asn Ala Ser Val Ser Ile Ala Leu Gly Thr Val Val Leu Ile
 65 70 75 80
 Ser Ala Leu Phe Thr Gly Glu Ile Leu Thr Asp Asp Ile Leu Ser Lys
 85 90 95
 Ile Gly Arg Asp Ser Arg Phe Leu Tyr Leu Met Gly Leu Thr Gly Arg
 100 105 110
 Leu Val Asn Asp Thr Lys Thr Tyr Gln Ala Glu Arg Gly Gln Gly Glu
 115 120 125
 Val Ala Ser Ala Val Gln Cys Tyr Met Lys Asp His Pro Glu Ile Ser
 130 135 140
 Glu Glu Glu Ala Leu Lys His Val Tyr Thr Ile Met Asp Asn Ala Leu
 145 150 155 160
 Asp Glu Leu Asn Arg Glu Phe Val Asn Asn Arg Asp Val Pro Asp Thr

165 170 175
 Cys Arg Arg Leu Val Phe Glu Thr Ala Arg Ile Met Gln Leu Phe Tyr
 180 185 190
 Met Asp Gly Asp Gly Leu Thr Leu Ser His Asn Met Glu Ile Lys Glu
 195 200 205
 His Val Lys Asn Cys Leu Phe Gln Pro Val Ala
 210 215

<210> 266
 <211> 423
 <212> PRT
 <213> Eucalyptus grandis

<400> 266
 Leu Asp Cys Glu Pro Val Val Gln Lys Pro Lys Leu Val Asp Pro Val
 1 5 10 15
 Val Gln Asp Ala Pro Lys Glu Lys Val Ile Glu Ala Val Pro Ser Ala
 20 25 30
 Met Pro Glu Glu Asp Glu Glu Ile Ile Lys Ser Val Val Glu Gly Lys
 35 40 45
 Met Pro Ser Tyr Ser Leu Glu Ser Lys Leu Gly Asp Cys Lys Arg Ala
 50 55 60
 Ala Ala Ile Arg Arg Glu Ala Leu Gln Arg Ile Thr Gly Lys Ser Leu
 65 70 75 80
 Ser Gly Leu Pro Leu Glu Gly Phe Asp Tyr Glu Ser Ile Leu Gly Gln
 85 90 95
 Cys Cys Glu Met Pro Val Gly Tyr Val Gln Ile Pro Val Gly Ile Ala
 100 105 110
 Gly Pro Leu Leu Leu Asp Gly Arg Glu Tyr Ser Val Pro Met Ala Thr
 115 120 125
 Thr Glu Gly Cys Leu Val Ala Ser Thr Asn Arg Gly Cys Lys Ala Ile
 130 135 140
 Phe Val Ser Gly Gly Ala Thr Ser Val Leu Leu Arg Asp Gly Met Thr
 145 150 155 160
 Arg Ala Pro Ile Val Arg Phe Gly Thr Ala Lys Arg Ala Ala Asp Leu
 165 170 175
 Lys Phe Phe Val Glu Asn Pro Ala Asn Phe Glu Ser Leu Ala Val Ile
 180 185 190
 Phe Asn Arg Ser Ser Arg Phe Ala Arg Leu Gln Ser Ile Lys Cys Ala
 195 200 205
 Ile Ala Gly Lys Asn Leu Tyr Met Arg Phe Ser Cys Ser Thr Gly Asp
 210 215 220
 Ala Met Gly Met Asn Met Val Ser Lys Gly Val Gln Asn Val Leu Asp
 225 230 235 240
 Phe Leu Gln Ser Asp Phe Pro Asp Met Asp Val Leu Gly Ile Ser Gly
 245 250 255
 Asn Phe Cys Ala Asp Lys Lys Pro Ala Ala Val Asn Trp Ile Glu Gly
 260 265 270
 Arg Gly Lys Ser Val Val Cys Glu Ala Thr Ile Lys Gly Asp Val Val
 275 280 285
 Arg Lys Val Leu Lys Thr Ser Val Glu Ala Leu Val Glu Leu Asn Met
 290 295 300
 Leu Lys Asn Leu Thr Gly Ser Ala Met Ala Gly Ala Leu Gly Gly Phe
 305 310 315 320
 Asn Ala His Ala Ser Asn Ile Val Ala Ala Ile Phe Ile Ala Thr Gly
 325 330 335
 Gln Asp Pro Ala Gln Asn Val Glu Ser Ser His Cys Ile Thr Met Met
 340 345 350
 Glu Ala Ile Asn Asp Gly Lys Asp Leu His Val Ser Val Thr Met Pro
 355 360 365
 Ser Ile Glu Val Gly Thr Val Gly Gly Gly Thr Gln Leu Ala Ser Gln

370 375 380
 Ser Ala Cys Leu Asn Leu Leu Gly Val Lys Gly Ala Asn Lys Glu Leu
 385 390 395 400
 Ala Gly Ala Asn Ser Arg Leu Leu Ala Thr Val Val Ser Gly Ala Val
 405 410 415
 Leu Ala Ala Glu Leu Ser Ser
 420

<210> 267
 <211> 112
 <212> PRT
 <213> Pinus radiata

<400> 267
 Met Ser Leu Ile Ser Ala Val Pro Leu Ala Ser Ser Cys Val Ser Lys
 1 5 10 15
 Ser Leu Ile Ser Ser Val Arg Glu His Lys Ala Leu Arg Arg Ala Ile
 20 25 30
 Ala Thr Leu Gln Met Ser Arg Pro Gly Lys Ser Val Ala Ala Ser Thr
 35 40 45
 Arg Met Ser Ser Ala Thr Ala Gly Ser Asp Asp Gly Val Lys Arg Arg
 50 55 60
 Ile Gly Asp Tyr His Ser Asn Leu Trp Glu Asp Asn Phe Ile Gln Ser
 65 70 75 80
 Leu Ser Ser Pro Tyr Gly Ala Ser Ser Tyr Gly Glu His Ala Asp Arg
 85 90 95
 Leu Ile Gly Glu Val Lys Gly Ile Phe Asn Ser Phe Ser Ile Ala Asp
 100 105 110

<210> 268
 <211> 165
 <212> PRT
 <213> Pinus radiata

<400> 268
 Met Ser Leu Ile Ser Ala Val Pro Leu Ala Ser Ser Ser Val Ser Lys
 1 5 10 15
 Ser Leu Ile Ser Ser Val Arg Glu His Lys Ala Leu Arg Arg Ala Ile
 20 25 30
 Ala Thr Leu Gln Met Ser Arg Pro Gly Lys Ser Val Ala Ala Ser Thr
 35 40 45
 Lys Met Ser Ser Ala Thr Ala Gly Ser Asp Asp Gly Val Lys Arg Arg
 50 55 60
 Ile Gly Asp Tyr His Ser Asn Leu Trp Asp Asp Asn Val Ile Gln Ser
 65 70 75 80
 Leu Ser Ser Pro Tyr Gly Ala Ser Ser Tyr Gly Glu His Ala Asp Arg
 85 90 95
 Leu Ile Gly Glu Val Lys Glu Ile Phe Asn Ser Phe Ser Ile Ala Asp
 100 105 110
 Gly Glu Leu Thr Ser Pro Val Asn Asp Leu Leu Gln Gln Leu Trp Met
 115 120 125
 Val Asp Asn Val Glu Arg Leu Gly Ile Asp Arg His Phe Gln Thr Glu
 130 135 140
 Ile Lys Val Ala Leu Asp Tyr Gly Tyr Arg Tyr Trp Ser Glu Lys Gly
 145 150 155 160
 Ile Glu Cys Gly Glu
 165

<210> 269
 <211> 144
 <212> PRT

<213> Pinus radiata

<400> 269

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Ser Thr Leu Gln Leu Ser Arg Arg Gly Lys Pro Val Thr Ala Cys Lys
 1           5           10           15
Lys Val Ser Leu Ser Thr Ala Val Ser Asp Asp Gly Ala Lys Arg Arg
          20           25           30
Val Gly Asp His His Ser Asn Leu Trp Asp Asp Asn Phe Ile Lys Ser
          35           40           45
Leu Ser Ser Pro Tyr Gly Ala Ser Ser Tyr Arg Glu His Ala Asp Arg
          50           55           60
Val Ile Gly Glu Val Lys Glu Ile Phe Asn Ser Leu Ser Met Thr Asp
65           70           75           80
Gly Glu Leu Ile Ser Pro Asp Asn Asp Leu Leu Gln Arg Leu Ser Met
          85           90           95
Val Asp Asn Ile Glu Arg Leu Gly Ile Asp Arg His Phe Gln Thr Glu
          100          105          110
Ile Lys Leu Thr Leu Asp Tyr Val Tyr Ser Tyr Trp Ser Glu Lys Gly
          115          120          125
Ile Gly Tyr Gly Arg Glu Ser Ala Ile Thr Asp Leu Asn Thr Thr Ser
130           135           140

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<210> 270

<211> 106

<212> PRT

<213> Pinus radiata

<400> 270

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Gly Thr Lys Ala Lys Gly Asn Lys Gln Leu Gln Asn Asn Val Ile Lys
 1           5           10           15
Val Ile Cys Asn Thr Asp Lys Ser Arg Gly Phe Asn Val Leu Arg Asp
          20           25           30
Val Ser Met Pro Gln Ile Met Ile Lys Ser Cys Lys Val Ser Pro Asp
          35           40           45
Ala Arg Pro Tyr Gln Asn Leu Gly Gly Pro Ala Ser Ser Glu Arg Pro
          50           55           60
Phe Leu Ala Phe Phe Ala Gly Gln Met His Gly Thr Leu Arg Pro Glu
65           70           75           80
Ile Leu Lys His Trp Gly Asn Glu Thr Asp Pro Asn Met Lys Ile Phe
          85           90           95
Ala Val Gly Gln Ser His Pro Gly Ser Leu
          100          105

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<210> 271

<211> 169

<212> PRT

<213> Eucalyptus grandis

<400> 271

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Lys Ala Arg Ala Val Trp Glu Asn Phe Lys Asp Asn Pro Leu Phe Asp
 1           5           10           15
Ile Ser Thr Asp His Pro Thr Thr Tyr Tyr Glu Asp Met Gln Arg Ala
          20           25           30
Val Phe Cys Leu Cys Pro Leu Gly Trp Ala Pro Trp Ser Pro Arg Leu
          35           40           45
Val Glu Ala Val Val Phe Gly Cys Ile Pro Val Ile Ile Ala Asp Asp
          50           55           60
Ile Val Leu Pro Phe Ala Asp Ala Ile Pro Trp Glu Glu Ile Gly Val
65           70           75           80
Phe Val Ala Glu Glu Asp Val Pro Ser Leu Asp Thr Ile Leu Thr Ser
          85           90           95

```

Ile Ser Pro Glu Val Ile Leu Arg Lys Gln Arg Leu Leu Ala Asn Pro
 100 105 110
 Ser Met Lys Arg Ala Met Leu Phe Pro Gln Pro Ala Gln Ser Gly Asp
 115 120 125
 Ala Phe His Gln Ile Leu Asn Gly Leu Ala Arg Lys Leu Pro His His
 130 135 140
 Arg Ser Val Tyr Leu Lys Pro Gly Glu Lys Val Leu Asn Trp Thr Ala
 145 150 155 160
 Gly Pro Val Gly Asp Leu Lys Pro Trp
 165

<210> 272

<211> 146

<212> PRT

<213> Eucalyptus grandis

<400> 272

Met Ser Gln Val Ser Ala Thr Pro Cys Ala Pro Pro Asn Lys Glu Thr
 1 5 10 15
 Gly His Val Ile Glu Arg Arg Ser Ala Gly Tyr His Pro Ser Val Trp
 20 25 30
 Gly Asp Tyr Phe Leu Lys Tyr Asp Ser Pro Ser Asn Ser Val Lys Phe
 35 40 45
 Lys Phe Leu Gly Arg Val Glu Gly Gln Ile Glu Glu Leu Lys Gly Glu
 50 55 60
 Val Lys Lys Met Leu Ile Asp Val Val Asp Lys Pro Leu Pro Lys Leu
 65 70 75 80
 His Leu Ile Asp Gln Ile Gln Arg Leu Gly Ile Glu Tyr His Phe Glu
 85 90 95
 Arg Glu Val Asp Glu Gln Leu Glu Gln Ile His Lys Ser Tyr Ser Arg
 100 105 110
 Leu Asp His Glu Asp Phe Lys Val Asp Asp Leu His Met Val Ala Leu
 115 120 125
 Ile Phe Arg Leu Leu Arg Gln His Gly Tyr Asn Ile Ser Ser Glu Ile
 130 135 140
 Phe Asp
 145

<210> 273

<211> 132

<212> PRT

<213> Eucalyptus grandis

<400> 273

Lys Lys Met Leu Ile Asp Ala Val Asp Lys Pro Leu Pro Lys Leu His
 1 5 10 15
 Leu Ile Asp Gln Ile Gln Arg Leu Gly Ile Glu Tyr His Phe Glu Arg
 20 25 30
 Glu Val Asp Glu Gln Leu Glu Gln Ile His Lys Ser Tyr Ser Arg Leu
 35 40 45
 Asp His Glu Asp Phe Lys Val Asp Asp Leu His Thr Val Ala Leu Ile
 50 55 60
 Phe Arg Leu Leu Arg Gln His Gly Tyr Asn Ile Ser Ser Glu Val Phe
 65 70 75 80
 Asp Lys Phe Lys Ile Ala Thr Gly Thr Ser Glu Ser Arg Leu Ile Ser
 85 90 95
 Asp Val Arg Gly Leu Leu Ser Leu Tyr Glu Ala Cys His Leu Arg Cys
 100 105 110
 His Gly Asp Ser Ile Leu Asp Glu Ala Leu Pro Phe Ala Thr Thr His
 115 120 125
 Leu Glu Ser Ile

130

<210> 274
 <211> 116
 <212> PRT
 <213> Eucalyptus grandis

<400> 274

Met	Ser	Gln	Val	Ser	Ala	Thr	Pro	Cys	Ala	Pro	Ser	Asn	Lys	Gly	Thr
1				5					10					15	
Gly	His	Val	Ile	Glu	Arg	Arg	Ser	Ala	Gly	Tyr	His	Pro	Ser	Val	Trp
		20					25					30			
Gly	Asp	Tyr	Phe	Leu	Lys	Tyr	Asp	Ser	Pro	Ser	Asn	Ser	Val	Lys	Phe
	35					40					45				
Lys	Phe	Leu	Gly	Arg	Val	Glu	Gly	Gln	Ile	Glu	Glu	Leu	Lys	Gly	Glu
	50				55					60					
Val	Lys	Lys	Met	Leu	Thr	Asp	Ile	Met	Asp	Lys	Pro	Leu	Gln	Lys	Leu
65				70					75					80	
His	Leu	Ile	Asp	Gln	Ile	Gln	Arg	Leu	Gly	Ile	Glu	Tyr	His	Phe	Glu
		85						90						95	
Arg	Glu	Ile	Asp	Glu	Gln	Leu	Glu	Gln	Ile	His	Lys	Ser	Tyr	Ser	Arg
	100						105						110		
Leu	Asp	His	Glu												
		115													

<210> 275
 <211> 214
 <212> PRT
 <213> Pinus radiata

<400> 275

Met	Ala	Thr	Phe	Ser	Asp	Glu	Thr	Pro	Val	Ser	Ser	Leu	Ala	Cys	Gly
1				5					10					15	
Leu	Ser	Ser	Asn	Ser	Gly	Leu	Ile	Arg	Arg	Thr	Ala	Asn	Pro	His	Pro
			20					25					30		
Asn	Val	Trp	Gly	Tyr	Glu	Phe	Val	Asn	Ser	Leu	Lys	Ser	Pro	Tyr	Ala
	35					40					45				
Asn	Ser	Ser	Tyr	Arg	Glu	Arg	Ala	Glu	Thr	Leu	Val	Ser	Glu	Ile	Lys
	50				55					60					
Ala	Met	Leu	Asn	Thr	Ala	Ile	Ala	Gly	Asp	Gly	Asp	Leu	Met	Ile	Thr
65				70					75					80	
Pro	Ser	Ala	Tyr	Asp	Thr	Ala	Trp	Ile	Ala	Arg	Val	Pro	Ala	Ile	Asp
			85					90					95		
Gly	Ser	Pro	Arg	Pro	Gln	Phe	Pro	Gln	Thr	Val	Asp	Trp	Ile	Leu	Lys
		100					105						110		
Asn	Gln	Leu	Lys	Asp	Gly	Ser	Trp	Gly	Thr	Gln	Ser	His	Phe	Leu	Leu
		115					120					125			
Ser	Asp	Arg	Leu	Leu	Ala	Thr	Leu	Ser	Cys	Val	Leu	Ala	Leu	Leu	Lys
	130					135					140				
Trp	Lys	Val	Gly	Asp	Ala	Gln	Val	Gln	Gln	Gly	Ile	Lys	Phe	Ile	Arg
145				150					155					160	
Ser	Asn	Leu	Leu	Lys	Asp	Glu	Ser	Asp	Glu	Asp	Ser	Leu	Val	Thr	Asp
			165					170					175		
Phe	Glu	Val	Asn	Phe	Pro	Phe	Leu	Leu	Arg	Glu	Ala	Gln	Ser	Phe	Gln
		180					185						190		
Leu	Glu	Leu	Pro	Tyr	Asp	Leu	Pro	Tyr	Ile	His	Lys	Leu	Gln	Met	Lys
		195					200						205		
Arg	Gln	Glu	Arg	Leu	Ala										
		210													

<210> 276

<211> 462

<212> PRT

<213> Pinus radiata

<400> 276

Arg Asp Ser Ala Phe Thr Asp Leu Asn Thr Thr Ala Leu Gly Phe Arg
 1 5 10 15
 Ile Phe Arg Leu His Gly Tyr Thr Val Ser Ser Asp Ala Phe Glu His
 20 25 30
 Phe Lys Asp Gln Met Gly Gln Phe Ser Ala Ser Ala Asn Asp Thr Glu
 35 40 45
 Leu Gln Ile Arg Ser Val Phe Asn Leu Phe Arg Ala Ser Leu Ile Ala
 50 55 60
 Phe Pro Glu Glu Lys Val Leu Glu Glu Ala Glu Asn Phe Ala Ala Ala
 65 70 75 80
 Tyr Leu Lys Ala Ala Leu Gln Thr Leu Pro Val Ser Gly Leu Ser Arg
 85 90 95
 Glu Ile Gln Tyr Val Phe Asp Tyr Arg Trp His Ser Asn Leu Pro Arg
 100 105 110
 Leu Glu Ala Arg Ser Tyr Val Asp Ile Leu Ala Asp Asn Thr Ile Ser
 115 120 125
 Gly Thr Pro Asp Ala Asn Thr Lys Lys Leu Leu Glu Leu Ala Lys Leu
 130 135 140
 Glu Phe Asn Ile Phe His Ser Leu Gln Gln Lys Glu Leu Gln Cys Leu
 145 150 155 160
 Trp Arg Trp Trp Lys Glu Trp Gly Cys Pro Glu Leu Thr Phe Val Arg
 165 170 175
 His Arg Tyr Val Glu Phe Tyr Thr Leu Val Ser Gly Thr Asp Met Val
 180 185 190
 Pro Glu His Ala Ala Phe Arg Leu Ser Phe Val Lys Thr Cys His Leu
 195 200 205
 Ile Thr Ile Leu Asp Asp Met Tyr Asp Thr Phe Gly Thr Ile Asp Glu
 210 215 220
 Leu Arg Leu Phe Thr Ala Ala Val Lys Arg Trp Asp Pro Ser Ala Thr
 225 230 235 240
 Glu Cys Leu Pro Glu Tyr Met Lys Gly Val Tyr Met Val Leu Tyr Glu
 245 250 255
 Thr Val Asn Glu Met Ala Lys Glu Ala Gln Lys Ser Gln Gly Arg Asp
 260 265 270
 Thr Leu Gly Tyr Val Arg Gln Ala Leu Glu Asp Tyr Ile Gly Ser Tyr
 275 280 285
 Leu Lys Glu Ala Glu Trp Ile Ala Thr Gly Tyr Val Pro Thr Phe Gln
 290 295 300
 Glu Tyr Phe Glu Asn Gly Lys Leu Ser Ser Gly His Arg Ile Ala Thr
 305 310 315 320
 Leu Gln Pro Ile Leu Thr Leu Ser Ile Pro Phe Pro His His Ile Leu
 325 330 335
 Gln Glu Ile Asp Phe Pro Ser Lys Phe Asn Asp Tyr Ala Cys Ser Ile
 340 345 350
 Leu Arg Leu Arg Gly Asp Thr Arg Cys Tyr Lys Ala Asp Ser Ala Arg
 355 360 365
 Gly Glu Glu Ala Ser Cys Ile Ser Cys Tyr Met Lys Glu Asn Pro Gly
 370 375 380
 Ser Thr Gln Glu Asp Ala Leu His His Ile Asn Gly Met Ile Glu Asp
 385 390 395 400
 Met Ile Lys Lys Leu Asn Trp Glu Phe Leu Lys Pro Asp Asn Asn Ala
 405 410 415
 Pro Ile Ser Ser Lys Lys Asn Ala Phe Asn Ile Ser Arg Gly Leu His
 420 425 430
 His Phe Tyr Asn Tyr Arg Asp Gly Tyr Ser Val Ala Ser Asn Glu Thr
 435 440 445

Lys Asp Leu Val Ile Lys Thr Val Leu Glu Pro Val Leu Met
 450 455 460

<210> 277
 <211> 98
 <212> PRT
 <213> Pinus radiata

<400> 277
 Leu Gly Glu Asp Ser Leu Thr Gly Thr Pro Asp Val Asn Thr Lys Lys
 1 5 10 15
 Leu Leu Glu Leu Ser Lys Leu Glu Phe Asn Ile Phe His Ser Val Gln
 20 25 30
 Gln Lys Glu Leu Gln Cys Leu Ser Arg Trp Trp Lys Glu Ser Gly Ser
 35 40 45
 Pro Glu Leu Thr Phe Ala Arg His Arg Tyr Val Glu Phe Tyr Thr Leu
 50 55 60
 Val Cys Gly Ile Asp Met Glu Pro Lys Asp Ala Ala Phe Arg Leu Ser
 65 70 75 80
 Phe Val Lys Met Cys His Leu Ile Thr Ile Leu Asp Asp Ile Tyr Asp
 85 90 95
 Thr Phe

<210> 278
 <211> 63
 <212> PRT
 <213> Pinus radiata

<400> 278
 Thr Glu Cys Leu Pro Glu Tyr Met Lys Gly Val Tyr Met Val Leu Tyr
 1 5 10 15
 Glu Thr Val Asn Glu Met Ala Lys Glu Ala Gln Lys Ser Gln Gly Arg
 20 25 30
 Asp Thr Leu Gly Tyr Val Arg Gln Ala Val Ile Thr Ile Asp Met Leu
 35 40 45
 Cys Ile Tyr Leu Asn Lys Gln Ile Leu Val Gly His Leu Phe Tyr
 50 55 60

<210> 279
 <211> 124
 <212> PRT
 <213> Pinus radiata

<400> 279
 Ala Asp Leu Leu Asp Glu Cys Gly Pro Leu Leu Lys Lys Ala His Ala
 1 5 10 15
 Phe Leu Glu Lys Ser Gln Val Gln Glu Asn Ser Pro Gly Glu Phe Ser
 20 25 30
 Thr Trp Tyr Arg His Ile Ser Lys Gly Ala Trp Thr Leu Ser Thr Arg
 35 40 45
 Asp His Gly Trp Val Val Ala Asp Cys Ser Ala Glu Gly Leu Lys Ala
 50 55 60
 Ala Leu Glu Leu Ser Gln Leu Pro Glu Asn Ile Val Gly Lys Pro Leu
 65 70 75 80
 Pro Gln Gln Arg Leu Phe Ala Cys Val Asn Tyr Leu Leu Ser Met Gln
 85 90 95
 Asn Thr Asp Gly Gly Tyr Ala Thr Tyr Asp Leu Thr Arg Ser Tyr Asn
 100 105 110
 Trp Leu Gly Thr Phe Asn Pro Ala Ala Ile Leu Gly
 115 120

<210> 280
 <211> 380
 <212> PRT
 <213> Eucalyptus grandis

<400> 280
 Met Asp Thr Asp Asn Lys Leu Phe Asn Val Gly Val Leu Leu Val Ala
 1 5 10 15
 Thr Leu Val Val Ala Lys Leu Ile Ser Ala Leu Leu Ile Pro Arg Ser
 20 25 30
 Gly Lys Arg Leu Pro Pro Val Val Arg Thr Trp Pro Val Val Gly Gly
 35 40 45
 Leu Leu Arg Phe Leu Lys Gly Pro Met Val Met Leu Arg Glu Glu Tyr
 50 55 60
 Pro Lys Leu Gly Ser Val Phe Thr Leu Asn Leu Leu Asn Lys Lys Ile
 65 70 75 80
 Thr Phe Phe Ile Gly Pro Glu Val Ser Ala His Phe Phe Lys Ala Ser
 85 90 95
 Glu Ser Asp Leu Ser Gln Gln Glu Val Tyr Gln Phe Asn Val Pro Thr
 100 105 110
 Phe Gly Pro Gly Val Val Phe Asp Val Asp Tyr Thr Ile Arg Gln Glu
 115 120 125
 Gln Phe Arg Phe Phe Thr Glu Ala Leu Arg Ile Asn Lys Leu Lys Gly
 130 135 140
 Tyr Val Asn Gln Met Val Met Glu Ala Glu Asp Tyr Phe Ser Lys Trp
 145 150 155 160
 Gly Asp Ser Gly Glu Val Asp Leu Lys Tyr Glu Leu Glu His Leu Thr
 165 170 175
 Ile Leu Thr Ala Ser Arg Cys Leu Leu Gly Arg Glu Val Arg Glu Lys
 180 185 190
 Leu Phe Asp Asp Val Ser Ala Leu Phe His Asp Leu Asp Asn Gly Met
 195 200 205
 Leu Pro Ile Ser Val Ile Phe Pro Tyr Leu Pro Ile Pro Ala His His
 210 215 220
 Arg Arg Asp Lys Ala Arg Lys Lys Leu Ser Glu Ile Phe Ala Asn Ile
 225 230 235 240
 Ile Ser Ser Arg Lys Cys Ala Gly Lys Ser Glu Glu Asp Met Leu Gln
 245 250 255
 Cys Phe Ile Asp Ser Lys Tyr Lys Asn Gly Arg Pro Thr Thr Glu Ala
 260 265 270
 Glu Val Thr Gly Leu Leu Ile Ala Ala Leu Phe Ala Gly Gln His Thr
 275 280 285
 Ser Ser Ile Thr Ser Val Trp Thr Gly Ala Tyr Leu Leu Thr Asn Lys
 290 295 300
 Lys Tyr Leu Ser Ala Val Ser Asn Glu Gln Lys His Leu Met Glu Lys
 305 310 315 320
 His Gly Asn Asn Val Asp His Asp Val Leu Ser Glu Met Asp Val Leu
 325 330 335
 Tyr Arg Ser Ile Lys Glu Ala Leu Arg Leu His Pro Pro Leu Ile Met
 340 345 350
 Leu Leu Arg Ser Ser His Ser Asp Phe Ser Val Lys Thr Arg Asp Gly
 355 360 365
 Lys Glu Tyr Glu Val Gly Glu Val Ser Val Leu Pro
 370 375 380

<210> 281
 <211> 177
 <212> PRT
 <213> Eucalyptus grandis

<400> 281

Met Trp Lys Leu Lys Ile Gly Glu Gly Ala Asn Asp Pro Tyr Leu Phe
 1 5 10 15
 Ser Leu Asn Asn Phe Val Gly Arg Gln Ile Trp Glu Phe Asp Pro Glu
 20 25 30
 Ala Gly Thr Pro Glu Glu Arg Ala Glu Val Glu Ala Ala Arg Gln Asn
 35 40 45
 Phe Tyr Asn Asn Arg Phe Lys Val Arg Pro Ser Ser Asp Leu Phe Trp
 50 55 60
 Arg Phe Gln Phe Leu Arg Glu Lys Asn Phe Lys Gln Thr Ile Pro Pro
 65 70 75 80
 Val Lys Ile Glu Asp Gly Glu Asp Ile Thr Tyr Glu Lys Ala Thr Ala
 85 90 95
 Ala Val Lys Arg Cys Val Ser Phe Trp Ser Thr Leu Gln Ser Ser His
 100 105 110
 Gly His Trp Pro Ala Glu Asn Ala Gly Pro Ile Ala Phe Tyr Phe Pro
 115 120 125
 Pro Leu Val Met Ser Leu Tyr Val Thr Gly His Leu Asn Asn Val Phe
 130 135 140
 His Ala Glu His Arg Arg Glu Ile Leu Arg Tyr Ile Tyr Tyr His Gln
 145 150 155 160
 Asn Glu Asp Gly Gly Trp Gly Leu His Ile Glu Gly His Ser Thr Met
 165 170 175
 Ile

<210> 282

<211> 91

<212> PRT

<213> Pinus radiata

<400> 282

His Ala Arg Gly Leu Gly Pro Pro Pro Ile Pro Val Asp Gln Phe Ser
 1 5 10 15
 Leu Ala Lys Leu Val Asp Ala Ile Gln Ile Met Leu Asn Pro Gln Val
 20 25 30
 Lys Asn Asn Ala Asp Ala Ile Ala Lys Ala Met Glu Asn Glu Asp Gly
 35 40 45
 Val Ser Gly Ala Val Lys Ala Phe His Lys His Leu Pro Lys Lys Met
 50 55 60
 Pro Gln Pro Leu Pro Pro Thr Asp His Ser Leu Ile Asp Ser Phe
 65 70 75 80
 Phe Thr Gly Val Gly Lys Val Phe Gly Cys Gly
 85 90

<210> 283

<211> 172

<212> PRT

<213> Pinus radiata

<400> 283

Trp Ile Glu Gly Arg Gly Lys Ser Val Val Cys Glu Ala Ile Ile Thr
 1 5 10 15
 Glu Ala Val Val Ser Lys Val Leu Lys Thr Thr Val Pro Ala Leu Leu
 20 25 30
 Glu Leu Asn Met Leu Lys Asn Leu Thr Gly Ser Ala Leu Ala Gly Ala
 35 40 45
 Met Gly Gly Phe Asn Ala His Ala Ser Asn Ile Val Ser Ala Val Phe
 50 55 60
 Ile Ala Thr Gly Gln Asp Pro Ala Gln Asn Ile Glu Ser Ser His Cys
 65 70 75 80

Ile Thr Met Met Glu Ala Ser Asn Asp Gly Lys Asp Leu His Val Ser
 85 90 95
 Val Thr Met Pro Cys Ile Glu Val Gly Thr Val Gly Gly Gly Thr Gln
 100 105 110
 Leu Ala Ser Gln Ala Ala Cys Leu Asn Met Leu Gly Val Lys Gly Ala
 115 120 125
 Asn Lys Glu Ser Pro Gly Ala Asn Ala Gln Thr Leu Ala Arg Ile Val
 130 135 140
 Ala Gly Ala Val Leu Ala Gly Glu Leu Ser Leu Met Ser Ala Leu Ala
 145 150 155 160
 Ala Gly Gln Leu Val Asn Ser His Met Lys Phe Asn
 165 170

<210> 284
 <211> 46
 <212> PRT
 <213> Pinus radiata

<400> 284
 Met Ala Thr Gly Gly Gly Ala Leu Asp Leu Ala Ser Gly Met Gly Gly
 1 5 10 15
 Asn Ile Glu Lys Glu Gln Met Leu Thr Ala Val Glu Glu Tyr Glu Lys
 20 25 30
 Tyr His Met Tyr Tyr Gly Gly Asp Glu Gly Ser Arg Lys Ser
 35 40 45

<210> 285
 <211> 137
 <212> PRT
 <213> Eucalyptus grandis

<400> 285
 Met Ser Lys Ala Gly Ala Met Asp Leu Ala Thr Gly Leu Gly Gly Lys
 1 5 10 15
 Met Asp Lys Ser Asp Val Leu Ser Ala Val Asp Lys Tyr Glu Lys Tyr
 20 25 30
 His Val Cys Tyr Gly Gly Asp Glu Glu Glu Arg Arg Ala Asn Tyr Ser
 35 40 45
 Asp Met Val Asn Lys Tyr Tyr Asp Leu Ala Thr Ser Phe Tyr Glu Phe
 50 55 60
 Gly Trp Gly Glu Ser Phe His Phe Ala His Arg Trp Lys Gly Glu Ser
 65 70 75 80
 Leu Arg Glu Ser Ile Lys Arg His Glu His Phe Leu Ala Leu Gln Leu
 85 90 95
 Gly Leu Lys Pro Gly His Lys Val Leu Asp Val Gly Cys Gly Ile Gly
 100 105 110
 Gly Pro Leu Arg Glu Ile Ala Arg Phe Ser Ser Ala Ser Val Thr Gly
 115 120 125
 Leu Asn Asn Asn Glu Tyr Gln Ile Thr
 130 135

<210> 286
 <211> 117
 <212> PRT
 <213> Pinus radiata

<400> 286
 Phe Arg Ile Trp Phe Asp Val Pro Val Val Leu Pro Pro Leu Thr Gln
 1 5 10 15
 Cys Phe Ala Asp Arg Ile Ser Leu Val Tyr Asp Pro His Thr Asp Glu
 20 25 30

Tyr Tyr Asn Ala Pro Gly Val Glu Thr Arg Val Pro Tyr Phe Gly Ser
 35 40 45
 Thr Glu Gly Met Lys Tyr Leu Asp Pro Cys Phe Lys Tyr Ile Thr Pro
 50 55 60
 Tyr Met Ser Ser Leu Val Lys Ser Leu Glu Asp Val Gly Tyr Val Asp
 65 70 75 80
 Gly Lys Ser Leu Phe Gly Ala Pro Tyr Asp Phe Arg Tyr Gly Pro Gly
 85 90 95
 Thr Lys Ser Ser Val Gly Ala Lys Tyr Leu Glu Asn Leu Arg Lys
 100 105 110
 Leu Val Glu Glu Ala
 115

<210> 287
 <211> 27
 <212> PRT
 <213> Eucalyptus grandis

<400> 287
 Gly Tyr Trp Asn Thr Met Asp Ile Ala His Asp Arg Ala Gly Phe Tyr
 1 5 10 15
 Ile Cys Trp Gly Cys Leu Val Trp Val Pro Ser
 20 25

<210> 288
 <211> 158
 <212> PRT
 <213> Pinus radiata

<400> 288
 Phe Ala Val Val Gly Pro Leu Gln Leu Thr Ser Tyr Pro Leu Ile Lys
 1 5 10 15
 Leu Val Gly Ile Arg Thr Gly Leu Pro Leu Pro Ser Leu Trp Glu Ile
 20 25 30
 Phe Ala Gln Leu Ala Val Tyr Phe Met Val Glu Asp Tyr Gly Asn Tyr
 35 40 45
 Trp Ile His Arg Trp Leu His Cys Lys Trp Gly Tyr Glu Lys Ile His
 50 55 60
 His Val His His Glu Phe Thr Ala Pro Met Gly Phe Ala Ala Pro Tyr
 65 70 75 80
 Ala His Trp Ser Glu Val Leu Ile Leu Gly Ile Pro Thr Phe Val Gly
 85 90 95
 Pro Ala Ile Ala Pro Gly His Met Ile Thr Phe Trp Cys Trp Val Val
 100 105 110
 Leu Arg Gln Val Glu Ala Ile Glu Thr His Ser Gly Tyr Asp Phe Pro
 115 120 125
 Trp Thr Leu Thr Lys Leu Ile Pro Phe Tyr Gly Gly Ala Glu Tyr His
 130 135 140
 Asp Tyr His His Tyr Val Gly Gly Gln Ser Gln Ser Asn Phe
 145 150 155

<210> 289
 <211> 113
 <212> PRT
 <213> Eucalyptus grandis

<400> 289
 Pro Ser Leu Trp Glu Ile Leu Ala Gln Leu Leu Val Tyr Phe Leu Ile
 1 5 10 15
 Glu Asp Tyr Thr Asn Tyr Trp Leu His Arg Leu Leu His Cys Lys Trp
 20 25 30

Gly Tyr Glu Lys Ile His Ser Val His His Glu Tyr Ser Ala Pro Ile
 35 40 45
 Gly Phe Ala Ala Pro Tyr Ala His Trp Ala Glu Val Leu Ile Leu Gly
 50 55 60
 Ile Pro Ser Phe Leu Gly Pro Ala Ile Val Pro Gly His Met Ile Thr
 65 70 75 80
 Leu Trp Leu Trp Ile Ala Leu Arg Gln Ile Glu Ala Ile Asp Tyr Ser
 85 90 95
 Gln Arg Val Arg Ile Ala Leu Glu Ser Tyr Glu Val His Ser Ile Leu
 100 105 110
 Trp

<210> 290
 <211> 128
 <212> PRT
 <213> Eucalyptus grandis

<400> 290
 Gly Tyr Gly Ser Met Val Gln Asn Cys Val Lys Ala Arg Ser Leu Leu
 1 5 10 15
 Ser Lys Leu Gly Ile Glu Val Thr Val Ala Asp Ala Arg Phe Cys Lys
 20 25 30
 Pro Leu Asp Ile Gly Leu Leu Arg Glu Leu Cys Glu Asn His Ala Phe
 35 40 45
 Leu Val Thr Val Glu Glu Gly Ser Ile Gly Gly Phe Gly Ser His Val
 50 55 60
 Ala Gln Phe Ile Ala Leu Asp Gly Arg Leu Asp Gly Arg Ile Lys Trp
 65 70 75 80
 Arg Pro Ile Val Leu Pro Asp Ala Tyr Val Glu His Thr Ser Pro Asn
 85 90 95
 Glu Gln Leu Ser Leu Ala Gly Leu Thr Gly His His Ile Ala Ala Thr
 100 105 110
 Val Leu Ser Leu Leu Gly Arg Thr Arg Glu Ala Leu Leu Leu Met Cys
 115 120 125

<210> 291
 <211> 109
 <212> PRT
 <213> Pinus radiata

<400> 291
 Met Ala Val Val Ser Ala Pro Gly Lys Val Leu Ile Thr Gly Ala
 1 5 10 15
 Tyr Leu Ile Leu Glu Lys Pro Asn Pro Gly Leu Val Leu Thr Thr Thr
 20 25 30
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 Leu Ala Lys Glu Ala Ile Tyr Lys Leu Ser Leu Lys Thr Leu Ser Leu
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 Thr Thr Thr Thr Val Ala Val Ser Pro Ala Phe Glu Gln Asp Arg Met
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<211> 148

<212> PRT

<213> Pinus radiata

<400> 293

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 65 70 75 80
 Val Glu Asp Thr Thr Met Asp Ala Val Gln Arg Arg Leu Met Phe Glu
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 Asp Glu Cys Ile Leu Val Asp Glu Glu Asp His Val Ile Gly His Asp
 100 105 110
 Ser Lys Tyr Asn Cys His Leu Met Glu Lys Ile Glu Ser Glu Asn Leu
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<210> 294

<211> 137

<212> PRT

<213> Eucalyptus grandis

<400> 294

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 35 40 45
 Cys Ile Leu Val Asp Glu Asn Asp Asn Val Val Gly His Glu Ser Lys
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85 90 95
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<212> PRT

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<400> 295

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 Lys Cys Ser Trp Leu Val Val Gln Ala Leu Glu Arg Ala Asn Glu Ser
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 65 70 75 80
 Val Ala Glu Val Lys Ala Val Tyr Arg Asp Leu Gly Ile Gln Asp Val
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 Phe Phe Glu Tyr Glu Arg Thr Ser Tyr Lys Glu Leu Ile Ser Ser Ile
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<210> 296

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<212> PRT

<213> Pinus radiata

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<212> PRT

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 Ser Ile Trp Asp Thr Phe Ser His Thr Pro Gly Lys Ile Ala Asp Gly
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 Ser Phe Xaa Xaa Gln Thr Pro Gly Lys Ile Val Asp Gly Ser Asn Gly
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 Asp Val Ala Val Asp Gln Tyr His Arg Tyr Lys Glu Asp Val Lys Leu
 65 70 75 80
 Ile Lys Asp Met Gly Val Asp Val Tyr Arg Phe Ser Ile Ser Trp Ser
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 Asp Arg Met Trp Leu
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<210> 302
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<400> 302
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 35 40 45
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 50 55 60
 Leu Glu His Ile Gln Arg Gln Ala Arg Lys Leu Gln Glu Gly Gly Trp
 65 70 75 80
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 35 40 45
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 50 55 60
 Ile Tyr Pro Leu Gly Arg Thr Arg Val Val Leu Lys Lys Ser Gly Asp
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 Gly Lys Pro Asp Ala Thr Phe Ser Phe Lys Asp Asp Asp Phe Ile Lys
 100 105 110
 Val Ala Thr Gly Lys Met Asn Pro Gln Ile Ala Phe Met Arg Gly Ala
 115 120 125
 Met Lys Ile Lys Gly Ser Leu Ser Ala Ala Gln Lys Phe Thr Pro Asp
 130 135 140
 Ile Phe Pro Lys Pro Ser Lys Met
 145 150

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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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(21) International Application Number: PCT/NZ99/00219 (22) International Filing Date: 16 December 1999 (16.12.99) (30) Priority Data: 09/215,504 17 December 1998 (17.12.98) US 60/146,441 29 July 1999 (29.07.99) US (71) Applicants (for all designated States except US): GENESIS RE-SEARCH AND DEVELOPMENT CORPORATION LIMITED [NZ/NZ]; 1 Fox Street, Parnell, Auckland (NZ). FLETCHER CHALLENGE FORESTS LIMITED [NZ/NZ]; 585 Great South Road, Penrose, Auckland (NZ). (72) Inventor; and (75) Inventor/Applicant (for US only): HAVUKKALA, Ilkka, Jaakko [FI/NZ]; 3/121 Atkin Avenue, Mission Bay, Auckland (NZ). (74) Agents: BENNETT, Michael, Roy et al.; West-Walker Bennett, Mobil on the Park, 157 Lambton Quay, Wellington (NZ).		(81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i> (88) Date of publication of the international search report: 3 August 2000 (03.08.00)
(54) Title: MATERIALS AND METHODS FOR THE MODIFICATION OF ISOPRENOID CONTENT, COMPOSITION AND METABOLISM		
(57) Abstract Novel isolated polynucleotides associated with plant isoprenoid biosynthetic pathways are provided, together with genetic constructs comprising such sequences. Methods for the modulation of the content, structure and metabolism of polypeptides involved in an isoprenoid biosynthetic pathway in target organisms are also disclosed, the methods comprising incorporating one or more of the polynucleotides or genetic constructs of the present invention into the genome of a target organism. Modulation of the content, structure and metabolism of such polypeptides produces modifications in the content, structure and metabolism of isoprenoids in the target organism.		

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CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
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DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

INTERNATIONAL SEARCH REPORT

International application No.

PCT/NZ99/00219

A. CLASSIFICATION OF SUBJECT MATTERInt. Cl. ⁷: C12N 5/10, 9/00, 15/29, 15/63


According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHEDMinimum documentation searched (classification system followed by classification symbols)
IPC7Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched
See Databases below.Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
Details in Supplemental Box V.**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	GenBank accession AJ011840 submitted 7 October 1998 by Clastre M.	1-29
X	GenBank accession AF019383 submitted 14 August 1997 by Lange BM et al.	1-29
X	GenBank accession Y15782 submitted 4 December 1997 by Camara B.	1-29
X	GenBank accession Y14333 submitted 28 July 1997 by Camara B.	1-29
X	GenBank accession AB003156 submitted 2 May 1997 by Suzuki H.	1-29

☒ Further documents are listed in the continuation of Box C
 ☐ See patent family annex

* Special categories of cited documents:	
"A" document defining the general state of the art which is not considered to be of particular relevance	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"E" earlier application or patent but published on or after the international filing date	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"O" document referring to an oral disclosure, use, exhibition or other means	"&" document member of the same patent family
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search
5 June 2000Date of mailing of the international search report
09 JUNE 2000Name and mailing address of the ISA/AU
AUSTRALIAN PATENT OFFICE
PO BOX 200, WODEN ACT 2606, AUSTRALIA
E-mail address: pct@ipaustalia.gov.au
Facsimile No. (02) 6285 3929Authorized officer

JULIE CAIRNDUFF
Telephone No : (02) 6283 2545

INTERNATIONAL SEARCH REPORT

International application No.

PCT/NZ99/00219

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	GenBank accession D78130 submitted 12 October 1995 by Sakakibara J et al.	1-29
X	GenBank accession AF061285 submitted 24 April 1998 by Back K et al.	1-29
X	GenBank accession U87908 submitted 31 January 1997 by Bohlmann J et al.	1-29
X	GenBank accession U87909 submitted 31 January 1997 by Bohlmann J et al.	1-29
X	GenBank accession AF006193 submitted 30 May 1997 by Bohlmann J et al.	1-29
X	GenBank accession U92266 submitted 5 March 1997 by Steele CL et al.	1-29
X	GenBank accession AF006195 submitted 30 May 1997 by Bohlmann J et al.	1-29
X	GenBank accession U60542 submitted 12 June 1996 by Kollipara KP et al.	1-29
X	GenBank accession L10390 submitted 22 September 1993 by Burnett RJ et al.	1-29
X	GenBank accession X54657 submitted 29 August 1990 by Chye ML.	1-29
X	GenBank accession U72146 submitted 21 September 1996 by Maldonado-Mendoza IE and Nessler CL.	1-29
X	GenBank accession X68652 submitted 7 October 1992 by Bach TJ.	1-29
X	GenBank accession X68651 submitted 7 October 1992 by Bach TJ.	1-29
X	GenBank accession X54659 submitted 29 August 1990 by Chye ML et al.	1-29
X	GenBank accession X15032 submitted 18 April 1989 by Caelles C.	1-29
X	GenBank accession M96068 submitted 27 April 1993 by Maldonado-Mendoza IE et al.	1-29
X	GenBank accession X96429 submitted 5 March 1996 by Chen XY et al.	1-29
X	GenBank accession U27535 submitted 23 May 1995 by Chen XY et al.	1-29
X	GenBank accession AB009029 submitted 20 November 1997 by Kushiro T.	1-29
X	GenBank accession AB009031 submitted 20 November 1997 by Kushiro T.	1-29
X	GenBank accession D89619 submitted 28 November 1996 by Shibuya M.	1-29
X	GenBank accession U02555 submitted 15 October 1993 by Matsuda SP.	1-29
X	GenBank accession U74319 submitted 15 October 1996 by Bak S et al.	1-29
X	GenBank accession Y09291 submitted 6 November 1996 by Weerck-Reichhart.	1-29

INTERNATIONAL SEARCH REPORT

International application No.
PCT/NZ99/00219

C (Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	GenBank accession AB014057 submitted 15 May 1998 by Kushiro T et al.	1-29
X	GenBank accession AB00930 submitted 20 November 1997 by Kushiro T.	1-29
X	GenBank accession Z83833 submitted 10 January 1997 by Warnecke D.	1-29
X	GenBank accession Z83832 submitted 10 January 1997 by Warnecke D.	1-29
X	GenBank accession U81312 submitted 7 December 1996 by Benveniste P.	1-29
X	GenBank accession U81313 submitted 7 December 1996 by Benveniste P.	1-29
X	GenBank accession AF045570 submitted 30 January 1998 by Tong Y and Nes WD.	1-29
X	GenBank accession U79669 submitted 25 November 1996 by Grebenok RJ et al.	1-29
X	GenBank accession U43683 submitted 20 December 1995 by Clouse JA.	1-29
X	GenBank accession U60205 submitted 6 June 1996 by Kaplan J and Li L.	1-29
X	GenBank accession U93162 submitted 11 March 1997 by Herrmann K.	1-29
X	GenBank accession D50559 submitted 15 May 1995 by Uwebe K.	1-29
X	GenBank accession U27099 submitted 12 May 1995 by Mandel MA et al.	1-29
X	GenBank accession Y14325 submitted 24 July 1997 by Cordier H.	1-29
X	GenBank accession U53706 submitted 6 April 1996 by Jeng CJ and Schweitzer ES.	1-29
X	GenBank accession U49260 submitted 15 February 1996 by Toth MJ et al.	1-29
X	GenBank accession Y17593 submitted 17 June 1998 by Cordier H.	1-29
X	GenBank accession Y09292 submitted 6 November 1996 by Werck-Reichhart D.	1-29
X	GenBank accession U50201 submitted 28 February 1996 by Poulton JE and Jurk S.	1-29
X	GenBank accession AF072736 submitted 16 June 1998 by Dharmawardhana D et al.	1-29
X	GenBank accession X56734 submitted 19 November 1990 by Hughes MA.	1-29
X	GenBank accession D83177 submitted 19 January 1996 by Inoue K.	1-29
X	GenBank accession U39228 submitted 23 October 1995 by Wiersma PA.	1-29
X	GenBank accession U26025 submitted 2 May 1995 by Zheng L and Poulton JE.	1-29

INTERNATIONAL SEARCH REPORT

International application No.

PCT/NZ99/00219

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	GenBank accession AB017026 submitted 20 August 1998 by Snider J et al.	1-29
X	GenPept accession CAA03409 submitted 21 August 1996 by Chenivresse X et al.	1-29
X	GenPept accession CAA76803 submitted 17 June 1998 by Cordier H.	1-29
X	AU, A 24637/99 (WASHINGTON STATE UNIVERSITY RESEARCH FOUNDATION) 21 January 1999 A01H 5/00, C07K 14/415, C12N 1/00, 5/04, 5/06, 9/00, 15/29, 15/52, 15/74, 15/79, 15/82, 15/84. See entire document.	1-29

INTERNATIONAL SEARCH REPORT

International application No.

PCT/NZ99/00219

Box I Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos :
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☒ Claims Nos : 1, part (11); 1, part (12); 2; 26, part (7); 26, part (8).
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
These claims refer to polynucleotide or polypeptide sequences comprising 40, 20 or 10 contiguous residues of sequences provided in SEQ. ID. NOs: 1-53, 78-286, 288-304. The scope of the claims encompasses many sequence fragments and it is not economically viable to search all possible combinations.
3. ☐ Claims Nos :
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a)

Box II Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

Continued in Supplemental Box

1. ☒ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/NZ99/00219

Supplemental Box I

(To be used when the space in any of Boxes I to VIII is not sufficient)

Continuation of Box No: II

The International Searching Authority has found that there are 37 separate inventions, wherein a single enzyme or protein type provides the special technical feature.

1. Nucleic and amino acid sequences SEQ. ID. NOs: 1, 252 encoding acetylcholinesterase precursor, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
2. Nucleic and amino acid sequences SEQ. ID. NOs: 2, 253 encoding deoxyxylulosephosphate synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
3. Nucleic and amino acid sequences SEQ. ID. NOs: 3, 4, 44, 254, 255, 295 encoding geranyltranstransferase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
4. Nucleic and amino acid sequences SEQ. ID. NOs: 5, 6, 256, 266 encoding farnesyltranstransferase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
5. Nucleic and amino acid sequences SEQ. ID. NOs: 7, 154, 258, 241 encoding squalene synthetase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
6. Nucleic and amino acid sequences SEQ. ID. NOs: 8-10, 155-157, 259-261, 242-244 encoding squalene monooxygenase. DNA probes or primers therefrom, transgenic cells and constructs containing the sequences. and methods of modulating biosynthesis of isoprenoid content and metabolism.
7. Nucleic and amino acid sequences SEQ. ID. NOs: 11, 82, 83, 262, 169, 170 encoding geranylgeranyl-diphosphate geranylgeranyltransferase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
8. Nucleic and amino acid sequences SEQ. ID. NOs: 12, 263 encoding trichodiene synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
9. Nucleic and amino acid sequences SEQ. ID. NOs: 13, 25-27, 84-88, 95, 115-118, 264, 276-278, 171-175, 182, 202-205 encoding pinene synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
10. Nucleic and amino acid sequences SEQ. ID. NOs: 14, 89, 90, 265, 176, 177 encoding abietadine synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.

Continued in Supplemental Box II

Supplemental Box II

(To be used when the space in any of Boxes I to VIII is not sufficient)

Continuation of Supplemental Box I

11. Nucleic and amino acid sequences SEQ. ID. NOs 15, 32, 91-94, 96-98, 131-135, 266, 283, 178-181, 183-185, 218-222 encoding hydroxymethylglutaryl-CoA reductase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
12. Nucleic and amino acid sequences SEQ. ID. NOs: 16-18, 99-102, 267-269, 186-189 encoding myrcene synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
13. Nucleic and amino acid sequences SEQ. ID. NOs: 19, 20, 26, 27, 103, 107, 108, 277, 278, 270, 271, 190, 194, 195 encoding limonene synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
14. Nucleic and amino acid sequences SEQ. ID. NOs: 21-23, 109-111, 272-274, 196-198 encoding cadinene synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
15. Nucleic and amino acid sequences SEQ. ID. NOs: 24, 114, 275, 201 encoding bisabolene synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
16. Nucleic and amino acid sequences SEQ. ID. NOs: 28, 119-122, 279, 206-209 encoding cycloartenol synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
17. Nucleic and amino acid sequences SEQ. ID. NOs: 29, 124-126, 280, 211-213 encoding obtusifolioside demethylase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
18. Nucleic and amino acid sequences SEQ. ID. NOs: 30, 281 encoding lupeol synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
19. Nucleic and amino acid sequences SEQ. ID. NOs: 31, 158, 159, 282, 245, 246 encoding udp-glucose:sterol glucosyltransferase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
20. Nucleic and amino acid sequences SEQ. ID. NOs: 33, 34, 160-162, 284, 285, 247-249 encoding sterolmethyltransferase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.

Continued in Supplemental Box III

Supplemental Box III

(To be used when the space in any of Boxes I to VIII is not sufficient)

Continuation of Supplemental Box II

21. Nucleic and amino acid sequences SEQ. ID. NOs: 35, 136, 286, 223 encoding lecithin:cholesterol acyl transferase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
22. Nucleic and amino acid sequences SEQ. ID. NOs: 36, 137, 287, 224 encoding sterol delta-7 reductase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
23. Nucleic and amino acid sequences SEQ. ID. NOs: 37, 38, 138-140, 288, 289, 225-227 encoding methyl sterol oxidase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
24. Nucleic and amino acid sequences SEQ. ID. NOs: 39, 290 encoding deoxyxylulosephosphate synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
25. Nucleic and amino acid sequences SEQ. ID. NOs: 40, 291 encoding phosphomevalonate kinase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
26. Nucleic and amino acid sequences SEQ. ID. NOs: 41, 50, 141, 142, 146, 292, 301, 228, 229, 233 encoding diphosphomevalonate decarboxylase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
27. Nucleic and amino acid sequences SEQ. ID. NOs: 42, 43, 143, 293, 294, 230 encoding isopentenyl-diphosphate delta-isomerase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
28. Nucleic and amino acid sequences SEQ. ID. NOs: 45, 296 encoding estradiol dehydrogenase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
29. Nucleic and amino acid sequences SEQ. ID. NOs: 46-49, 144, 145, 297-300, 231-232 encoding furostanol glucosidase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
30. Nucleic and amino acid sequences SEQ. ID. NOs: 51, 52, 147-153, 302, 303, 234-240 encoding oxysterol-binding protein, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.

Continued in Supplemental Box IV

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Supplemental Box IV

(To be used when the space in any of Boxes I to VIII is not sufficient)

Continuation of Supplemental Box III

31. Nucleic and amino acid sequences SEQ. ID. NOs: 53, 304 encoding sterol carrier protein. DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
32. Nucleic and amino acid sequences SEQ. ID. NOs: 78, 79, 127-130, 165, 166, 214-217 encoding sterol 14-demethylase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
33. Nucleic and amino acid sequences SEQ. ID. NOs: 82, 83, 169, 170 encoding geranylgeranyl diphosphate, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
34. Nucleic and amino acid sequences SEQ. ID. NOs: 104-106, 164, 191-193, 251 encoding CXPS/transketolase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
35. Nucleic and amino acid sequences SEQ. ID. NOs: 112, 113, 199, 200 encoding sabinene synthase. DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
36. Nucleic and amino acid sequences SEQ. ID. NOs: 123, 210 encoding beta-amyrin synthase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.
37. Nucleic and amino acid sequences SEQ. ID. NOs: 163, 250 encoding sterol desaturase, DNA probes or primers therefrom, transgenic cells and constructs containing the sequences, and methods of modulating biosynthesis of isoprenoid content and metabolism.

The above inventions have been allocated into the following groups for searching purposes:

- A: Inventions 1 to 6.
- B: Inventions 7 to 10.
- C: Inventions 11 and 12.
- D: Inventions 13 to 15.
- E: Inventions 16 to 20.
- F: Inventions 21 to 26.
- G: Inventions 27 to 30.
- H: Inventions 31 to 37.

INTERNATIONAL SEARCH REPORT

International application No.

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Supplemental Box V

(To be used when the space in any of Boxes I to VIII is not sufficient)

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used):

GenBank, EMBL, PDB Nucleic Acids, SWISS-PROT, GenPept, PIR, TREMBL - SEQ. ID. NOS: 1-53, 78-286, 288-304

WPIDS: Keywords used - acetylcholinesterase precursor, deoxyxylulosephosphate synthase, dxps, geranyltranstransferase, farnesyl diphosphate synthase, farnesyltranstransferase, farnesyl diphosphate farnesyltransferase, presqualene diphosphate, squalene synthetase, squalene monooxygenase, squalene epoxidase, geranylgeranyl diphosphate geranylgeranyltransferase, prephytoene diphosphate synthase, trichodiene synthase, pinene synthase, abietadine synthase, hydroxymethylglutaryl coa reductase, myrcene synthase, limonene synthase, cadinene synthase, bisabolene synthase, cycloartenol synthase, epoxysqualene cycloarteno cyclase, obtusifolioside demethylase, lupeol synthase, udp glucose sterol glucosyl transferase, sterol glucosyltransferase, sterolmethyltransferase, lecithin cholesterol acyl transferase, phospholipid cholesterol acyltransferase, sterol delta 7 reductase, methyl sterol oxidase, deoxyxylulosephosphate synthase, dxps, diphosphomevalonate decarboxylase, phosphomevalonate kinase, isopentenyl diphosphate delta isomerase, estradioldehydrogenase, furostanol glucosidase, oxysterol binding protein, sterol carrier protein, sterol 14 demethylase, sesquiterpene cyclase, trichodiene synthase, dxps transketolase, sabinene synthase, beta amyrin synthase, sterol desaturase, pinus radiata, p radiata, pine, or pinus, eucalyptus grandis, e grandis, eucalyptus, isoprenylation, isoprenoid

INTERNATIONAL SEARCH REPORT

This Annex lists the known "A" publication level patent family members relating to the patent documents cited in the above-mentioned international search report. The Australian Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

Patent Document Cited in Search Report		Patent Family Member	
AU	24637/99	WO	9937139
END OF ANNEX			